

High diversity of *Diaporthe* species associated with dieback diseases in China, with twelve new species described

Qin Yang¹, Xin-Lei Fan¹, Vladimiro Guarnaccia^{2,3}, Cheng-Ming Tian¹,

1 *The Key Laboratory for Silviculture and Conservation of the Ministry of Education, Beijing Forestry University, Beijing 100083, P.R. China, V. Guarnaccia* **2** *Westerdijk Fungal Biodiversity Institute, Uppsalalaan 8, 3584 CT, Utrecht, The Netherlands* **3** *Department of Plant Pathology, University of Stellenbosch, Matieland 7602, South Africa*

Corresponding author: Cheng-Ming Tian (chengmt@bjfu.edu.cn)

Academic editor: George Mugambi | Received 24 May 2018 | Accepted 3 September 2018 | Published 17 September 2018

Citation: Yang Q, Fan X-L, Guarnaccia V, Tian C-M (2018) High diversity of *Diaporthe* species associated with dieback diseases in China, with twelve new species described. MycoKeys 39: 97–149. <https://doi.org/10.3897/mycokeys.39.26914>

Abstract

Diaporthe species have often been reported as important plant pathogens, saprobes and endophytes on a wide range of plant hosts. Although several *Diaporthe* species have been recorded in China, little is known about species able to infect forest trees. Therefore, extensive surveys were recently conducted in Beijing, Heilongjiang, Jiangsu, Jiangxi, Shaanxi and Zhejiang Provinces. The current results emphasised on 15 species from 42 representative isolates involving 16 host genera using comparisons of DNA sequence data for the nuclear ribosomal internal transcribed spacer (ITS), *calmodulin* (*cal*), histone H3 (*his3*), partial translation elongation factor-1 α (*tef1*) and β -tubulin (*tub2*) gene regions, as well as their morphological features. Three known species, *D. biguttulata*, *D. eres* and *D. unshiuensis*, were identified. In addition, twelve novel taxa were collected and are described as *D. acerigena*, *D. alangii*, *D. betulina*, *D. caryae*, *D. cercidis*, *D. chensiensis*, *D. cinnamomi*, *D. conica*, *D. fraxinicola*, *D. kadsurae*, *D. padina* and *D. ukurunduensis*. The current study improves the understanding of species causing diebacks on ecological and economic forest trees and provides useful information for the effective disease management of these hosts in China.

Keywords

Dieback, DNA phylogeny, Systematics, Taxonomy

Introduction

The genus *Diaporthe* Nitschke represents a cosmopolitan group of fungi occupying diverse ecological behaviour as plant pathogens, endophytes and saprobes (Muralli et al. 2006, Rossman et al. 2007, Garcia-Reyne et al. 2011, Udayanga et al. 2011, 2012a, b, 2014a, b, 2015, Gomes et al. 2013, Fan et al. 2015, Du et al. 2016, Dissanayake et al. 2017b, Guarnaccia and Crous 2017, Yang et al. 2017a, b, 2018, Guarnaccia et al. 2018, Marin-Felix et al. 2018). *Diaporthe* species are responsible for diseases on a wide range of plant hosts, including agricultural crops, forest trees and ornamentals, some of which are economically important. Several symptoms such as root and fruit rots, dieback, stem cankers, leaf spots, leaf and pod blights and seed decay are caused by *Diaporthe* spp. (Uecker 1988, Rehner and Uecker 1994, Mostert et al. 2001, Santos et al. 2011, Thompson et al. 2011, Udayanga et al. 2011). For example, *D. ampelina*, the causal agent of Phomopsis cane and leaf spot, is known as a severe pathogen of grapevines (Hewitt and Pearson 1988), infecting all green tissues and causing yield reductions of up to 30% in temperate regions (Erincik et al. 2001). *Diaporthe citri* is another well-known pathogen exclusively found on *Citrus* spp. causing melanose, stem-end rot and gummosis in all the citrus production areas except Europe (Mondal et al. 2007, Udayanga et al. 2014a, Guarnaccia and Crous 2017, 2018). Similarly, stem canker, attributed to several *Diaporthe* spp., is one of the most important diseases of sunflower (*Helianthus annuus*) worldwide (Muntañola-Cvetković et al. 1981, Thompson et al. 2011).

Several species of *Diaporthe* include a broad number of endophytes associated with hosts present in temperate and tropical regions (Udayanga et al. 2011). Gomes et al. (2013) considered that *D. endophytica* is a sterile endophyte on *Schinus terebinthifolius* and *Maytenus ilicifolia* based on molecular phylogeny. Huang et al. (2015) distinguished seven undescribed *Diaporthe* species associated with citrus in China. Moreover, some endophytes have been shown to act as opportunistic plant pathogens. For instance, *D. foeniculina* has been found as both endophyte and opportunistic pathogen on various herbaceous weeds, ornamentals and fruit trees (Udayanga et al. 2014a, Guarnaccia et al. 2016).

The genus *Diaporthe* (syn. *Phomopsis*) was established by Nitschke (1870). Species identification criteria in *Diaporthe* were originally based on host association, morphology and culture characteristics (Mostert et al. 2001, Santos and Phillips 2009, Udayanga et al. 2012). As a consequence, a broad increase in the number of proposed *Diaporthe* species occurred. More than 1000 epithets for *Diaporthe* and 950 for *Phomopsis* were listed in Index Fungorum (2018) (<http://www.indexfungorum.org/>) (accessed 1 March 2018). The abolishment of the dual nomenclature system for pleomorphic fungi raised the question about which generic name to use. Given that both names are well known amongst plant pathologists and have been equally used, Rossman et al. (2015) proposed that the name *Diaporthe* (Nitschke 1870) has priority over *Phomopsis* (Saccardo and Roumeguère 1884) and has been adopted as

the generic name in recent major studies (Gomes et al. 2013, Udayanga et al. 2014a, b, 2015, Fan et al. 2015, Huang et al. 2015, Du et al. 2016, Gao et al. 2017, Yang et al. 2017a, b, c, 2018).

The sexual morph of *Diaporthe* is characterised by immersed ascomata and an erumpent pseudostroma with elongated perithecial necks. Asci are unitunicate, clavate to cylindrical. Ascospores are fusoid, ellipsoid to cylindrical, hyaline, biseriate to uniseriate in the ascus and sometimes with appendages (Udayanga et al. 2011). The asexual morph is characterised by ostiolate conidiomata, with cylindrical phialides producing three types of hyaline, aseptate conidia (Udayanga et al. 2011). Previously, species identification of *Diaporthe* was largely referred to the assumption of host-specificity, leading to the proliferation of names (Gomes et al. 2013). More than one species of *Diaporthe* can colonise a single host, while one species can be associated with different hosts (Santos and Phillips 2009, Diogo et al. 2010, Santos et al. 2011, Gomes et al. 2013). In addition, considerable variability of the phenotype characters is present within a species (Rehner and Uecker 1994, Mostert et al. 2001, Santos et al. 2010, Udayanga et al. 2011, 2012a). Species identification is essential for understanding the epidemiology and plant diseases management and to guide the implementation of phytosanitary measures (Santos and Phillips 2009, Udayanga et al. 2011, Santos et al. 2017). Thus, molecular data are necessary to resolve *Diaporthe* taxonomy and, during the recent years, many species have been described through a polyphasic approach together with morphology (Gomes et al. 2013, Udayanga et al. 2014a, b, 2015, Huang et al. 2015, Gao et al. 2017, Guarnaccia and Crous 2017, Yang et al. 2018). Santos et al. (2017) revealed that the use of a five-loci dataset (ITS-*cal-his3-tef1-tub2*) is the optimal combination for species delimitation, showing the ribosomal ITS locus as the least informative, which is contrary to the result of Santos et al. (2010).

Although the classification of *Diaporthe* has been on-going, species are currently being identified based on a combination of morphological, cultural, phytopathological and phylogenetical analyses (Gomes et al. 2013, Huang et al. 2013, 2015, Udayanga et al. 2014a, b, 2015, Fan et al. 2015, Du et al. 2016, Gao et al. 2016, 2017, Guarnaccia and Crous 2017, Hyde et al. 2017, 2018, Guarnaccia et al. 2018, Jayawardena et al. 2018, Perera et al. 2018a, b, Tibpromma et al. 2018, Wanasinghe et al. 2018). However, fungi isolated from forest trees in China were recorded in old fungal literature without any living culture and molecular data (Teng 1963, Tai 1979, Wei 1979). The current study aimed to investigate the major ecological or economic trees in China by large-scale sampling and to identify isolates via morphology and multi-locus phylogeny based on modern taxonomic concepts. From 2015 to 2017, several surveys were conducted in six Provinces representing 16 host genera. The objectives of the present study were (i) to provide a multi-gene phylogeny for the genus *Diaporthe* based on a large set of freshly collected specimens in China; (ii) to identify *Diaporthe* taxa associated with disease symptoms or non-symptomatic tissues of various host genera distributed over six Provinces in China; (iii) to define the species limits of *D. eres* and closely related species based on multi-gene genealogies.

Table 1. Isolates and GenBank accession numbers used in the phylogenetic analyses of *Diaporthe*.

Species	Isolate	Host	Location	GenBank accession numbers				
				ITS	cal	his3	tef1	tub2
<i>D. acaciarium</i>	CBS 138862	<i>Acacia tortilis</i>	Tanzania	KP004460	N/A ^a	N/A ^a	N/A ^a	KP004509
<i>D. acaciigena</i>	CBS 129521	<i>Acacia retinodes</i>	Australia	KC343005	KC343247	KC343489	KC343731	KC343973
<i>D. acericola</i>	MFLUCC 17-0956	<i>Acer negundo</i>	Italy	KY964224	KY964137	N/A ^a	KY964180	KY964074
<i>D. acerigena</i>	CFCC 52554	<i>Acer tataricum</i>	China	MH121489	MH121413	MH121449	MH121531	N/A ^a
	CFCC 52555	<i>Acer tataricum</i>	China	MH121490	MH121414	MH121450	MH121532	N/A ^a
<i>D. acutispora</i>	CGMCC 3.18285	<i>Coffea</i> sp.	China	KX986764	KX999274	N/A ^a	KX999155	KX999195
<i>D. alangii</i>	CFCC 52556	<i>Alangium kurzii</i>	China	MH121491	MH121415	MH121451	MH121533	MH121573
	CFCC 52557	<i>Alangium kurzii</i>	China	MH121492	MH121416	MH121452	MH121534	MH121574
	CFCC 52558	<i>Alangium kurzii</i>	China	MH121493	MH121417	MH121453	MH121535	MH121575
	CFCC 52559	<i>Alangium kurzii</i>	China	MH121494	MH121418	MH121454	MH121536	MH121576
<i>D. alleghaniensis</i>	CBS 495.72	<i>Betula alleghaniensis</i>	Canada	KC343007	KC343249	KC343491	KC343733	KC343975
<i>D. alnea</i>	CBS 146.46	<i>Alnus</i> sp.	Netherlands	KC343008	KC343250	KC343492	KC343734	KC343976
<i>D. ambigua</i>	CBS 114015	<i>Pyrus communis</i>	South Africa	KC343010	KC343252	KC343494	KC343736	KC343978
<i>D. ampelina</i>	STEU2660	<i>Vitis vinifera</i>	France	AF230751	AY745026	N/A ^a	AY745056	JX275452
<i>D. amygdali</i>	CBS 126679	<i>Prunus dulcis</i>	Portugal	KC343022	KC343264	KC343506	AY343748	KC343990
<i>D. anacardii</i>	CBS 720.97	<i>Anacardium occidentale</i>	East Africa	KC343024	KC343266	KC343508	KC343750	KC343992
<i>D. angelicae</i>	CBS 111592	<i>Heracleum sphondylium</i>	Austria	KC343027	KC343269	KC343511	KC343753	KC343995
<i>D. apiculatum</i>	CGMCC 3.17533	<i>Camellia sinensis</i>	China	KP267896	N/A ^a	N/A ^a	KP267970	KP293476
<i>D. aquatica</i>	IFRDCC 3051	<i>Aquatic habitat</i>	China	JQ797437	N/A ^a	N/A ^a	N/A ^a	N/A ^a
<i>D. arctii</i>	CBS 139280	<i>Arctium lappa</i>	Austria	KJ590736	KJ612133	KJ659218	KJ590776	KJ610891
<i>D. arecae</i>	CBS 161.64	<i>Areca catechu</i>	India	KC343032	KC343274	KC343516	KC343758	KC344000
<i>D. arengae</i>	CBS 114979	<i>Arenga enngleri</i>	Hong Kong	KC343034	KC343276	KC343518	KC343760	KC344002
<i>D. aseana</i>	MFLUCC 12-0299a	Unknown dead leaf	Thailand	KT459414	KT459464	N/A ^a	KT459448	KT459432
<i>D. asheicola</i>	CBS 136967	<i>Vaccinium ashei</i>	Chile	KJ160562	KJ160542	N/Aa	KJ160594	KJ160518
<i>D. aspalathi</i>	CBS 117169	<i>Aspalathus linearis</i>	South Africa	KC343036	KC343278	KC343520	KC343762	KC344004
<i>D. australafricana</i>	CBS 111886	<i>Vitis vinifera</i>	Australia	KC343038	KC343280	KC343522	KC343764	KC344006
<i>D. baccae</i>	CBS 136972	<i>Vaccinium corymbosum</i>	Italy	KJ160565	N/A ^a	MF418264	KJ160597	N/A ^a

Species	Isolate	Host	Location	GenBank accession numbers				
				ITS	cal	bis3	ref1	tub2
<i>D. batatas</i>	CBS 122.21	<i>Ipomoea batatas</i>	USA	KC343040	KC343282	N/A ^a	KC343766	KC344008
<i>D. beilharziae</i>	BRIP 54792	<i>Indigofera australis</i>	Australia	JX862529	N/A ^a	N/A ^a	JX862535	KF170921
<i>D. benedicti</i>	BPI 893190	<i>Salix</i> sp.	USA	KM669929	KM669862	N/A ^a	KM669785	N/A ^a
<i>D. betulae</i>	CFCC 50469	<i>Betula platyphylla</i>	China	KT732950	KT732997	KT732999	KT733016	KT733020
	CFCC 50470	<i>Betula platyphylla</i>	China	KT732951	KT732998	KT733000	KT733017	KT733021
<i>D. betulicola</i>	CFCC 51128	<i>Betula albo-sinensis</i>	China	KX024653	KX024659	KX024661	KX024655	KX024657
	CFCC 51129	<i>Betula albo-sinensis</i>	China	KX024654	KX024660	KX024662	KX024656	KX024658
	CFCC 52560	<i>Betula albo-sinensis</i>	China	MH121495	MH121419	MH121455	MH121537	MH121577
<i>D. betulina</i>	CFCC 52561	<i>Betula costata</i>	China	MH121496	MH121420	MH121456	MH121538	MH121578
	CFCC 52562	<i>Betula platyphylla</i>	China	MH121497	MH121421	MH121457	MH121539	MH121579
	CBS 121004	<i>Juglans</i> sp.	USA	KC343134	KC343376	KC343618	KC343860	KC344102
<i>D. biconispora</i>	CGMCC 3.17252	<i>Citrus grandis</i>	China	KJ490597	KJ490539	KJ490539	KJ490476	KJ490418
<i>D. biguttulata</i>	CGMCC 3.17248	<i>Citrus limon</i>	China	KJ490582	N/A ^a	KJ490524	KJ490461	KJ490403
	CFCC 52584	<i>Juglans regia</i>	China	MH121519	MH121437	MH121477	MH121561	MH121598
	CFCC 52585	<i>Juglans regia</i>	China	MH121520	MH121438	MH121478	MH121562	MH121599
<i>D. biguttusis</i>	CGMCC 3.17081	<i>Libocarpus glabra</i>	China	KF576282	N/A ^a	N/A ^a	KF576257	KF576306
<i>D. bohemiae</i>	CPC 28222	<i>Vitis vinifera</i>	Czech Republic	MG281015	MG281710	MG281361	MG281536	MG281188
<i>D. brasiliensis</i>	CBS 133183	<i>Aspidosperma tomentosum</i>	Brazil	KC343042	KC343284	KC343526	KC343768	KC344010
	CBS 141542	<i>Tacinga inamoena</i>	Brazil	KY085927	N/A ^a	N/A ^a	KY115603	KY115600
<i>D. caatingaensis</i>	CFCC 51632	<i>Camptotheca acuminata</i>	China	KY203726	KY228877	KY228881	KY228887	KY228893
<i>D. canthii</i>	CBS 132533	<i>Canthium inerme</i>	South Africa	JX069864	KC843174	N/A ^a	KC843120	KC843230
<i>D. caryae</i>	CFCC 52563	<i>Carya illinoensis</i>	China	MH121498	MH121422	MH121458	MH121540	MH121580
	CFCC 52564	<i>Carya illinoensis</i>	China	MH121499	MH121423	MH121459	MH121541	MH121581
<i>D. cassines</i>	CPC 21916	<i>Cassine peragua</i>	South Africa	KF777155	N/A ^a	N/A ^a	KF777244	N/A ^a
<i>D. caulivora</i>	CBS 127268	<i>Glycine max</i>	Croatia	KC343045	KC343287	N/A ^a	KC343771	KC344013
<i>D. celeris</i>	CPC 28262	<i>Vitis vinifera</i>	Czech Republic	MG281017	MG281712	MG281363	MG281538	MG281190
<i>D. celastrina</i>	CBS 139.27	<i>Celastrus</i> sp.	USA	KC343047	KC343289	KC343531	KC343773	KC344015
	CFCC 52565	<i>Cercis chinensis</i>	China	MH121500	MH121424	MH121460	MH121542	MH121582
	CFCC 52566	<i>Cercis chinensis</i>	China	MH121501	MH121425	MH121461	MH121543	MH121583

Species	Isolate	Host	Location	GenBank accession numbers				
				ITS	cal	his3	ref1	tub2
<i>D. chamaeropsis</i>	CBS 454.81	<i>Chamaerops humilis</i>	Greece	KC343048	KC343290	KC343532	KC343774	KC344016
<i>D. charlesworthii</i>	BRIP 54884m	<i>Rapistrum rugostrum</i>	Australia	KJ197288	N/A ^a	N/A ^a	KJ197250	KJ197268
<i>D. chensiensis</i>	CFCC 52567	<i>Abies chensiensis</i>	China	MH121502	MH121426	MH121462	MH121544	MH121584
	CFCC 52568	<i>Abies chensiensis</i>	China	MH121503	MH121427	MH121463	MH121545	MH121585
<i>D. cichorii</i>	MFLUCC 17-1023	<i>Cichorium intybus</i>	Italy	KY964220	KY964133	N/A ^a	KY964176	KY964104
<i>D. cinnamomi</i>	CFCC 52569	<i>Cinnamomum</i> sp.	China	MH121504	N/A ^a	MH121464	MH121546	MH121586
	CFCC 52570	<i>Cinnamomum</i> sp.	China	MH121505	N/A ^a	MH121465	MH121547	MH121587
<i>D. cissampeli</i>	CBS 141331	<i>Cissampelos capensis</i>	South Africa	KX228273	N/A ^a	KX228366	N/A ^a	KX228384
<i>D. citri</i>	AR 3405	<i>Citrus</i> sp.	USA	KC843311	KC843157	N/A ^a	KC843071	KC843187
<i>D. citriasiatica</i>	CGMCC 3.15224	<i>Citrus unshiu</i>	China	JQ954645	KC357491	KJ490515	JQ954663	KC357459
<i>D. citrichinensis</i>	CGMCC 3.15225	<i>Citrus</i> sp.	China	JQ954648	KC357494	N/A ^a	JQ954666	N/A ^a
<i>D. collariana</i>	MFLU 17-2770	<i>Magnolia champaca</i>	Thailand	MG806115	MG783042	N/A ^a	MG783040	MG783041
<i>D. compacta</i>	CGMCC 3.17536	<i>Camellia sinensis</i>	China	KP267854	N/A ^a	KP293508	KP267928	KP293434
<i>D. conica</i>	CFCC 52571	<i>Alangium chinense</i>	China	MH121506	MH121428	MH121466	MH121548	MH121588
	CFCC 52572	<i>Alangium chinense</i>	China	MH121507	MH121429	MH121467	MH121549	MH121589
	CFCC 52573	<i>Alangium chinense</i>	China	MH121508	MH121430	MH121468	MH121550	MH121590
	CFCC 52574	<i>Alangium chinense</i>	China	MH121509	MH121431	MH121469	MH121551	MH121591
<i>D. convolvuli</i>	CBS 124654	<i>Convolvulus arvensis</i>	Turkey	KC343054	KC343296	KC343538	KC343780	KC344022
<i>D. crotalariae</i>	CBS 162.33	<i>Crotalaria spectabilis</i>	USA	KC343056	KC343298	KC343540	KC343782	KC344024
<i>D. cucurbitae</i>	CBS 136.25	<i>Arctium</i> sp.	Unknown	KC343031	KC343273	KC343515	KC343757	KC343999
<i>D. cuppatea</i>	CBS 117499	<i>Aspalathus linearis</i>	South Africa	KC343057	KC343299	KC343541	KC343783	KC344025
<i>D. cynaroidis</i>	CBS 122676	<i>Protea cynaroides</i>	South Africa	KC343058	KC343300	KC343542	KC343784	KC344026
<i>D. cytosporella</i>	FAU461	<i>Citrus limon</i>	Italy	KC843307	KC843141	N/A ^a	KC843116	KC843221
<i>D. diospyricola</i>	CPC 21169	<i>Diospyros wbyteaana</i>	South Africa	KF777156	N/A ^a	N/A ^a	N/A ^a	N/A ^a
<i>D. discoidispora</i>	ZJUD89	<i>Citrus unshiu</i>	China	KJ490624	N/A ^a	KJ490566	KJ490503	KJ490445
<i>D. dorycnii</i>	MFLUCC 17-1015	<i>Dorycnium hirsutum</i>	Italy	KY964215	N/A ^a	N/A ^a	KY964171	KY964099
<i>D. elaeagni-glabrae</i>	CGMCC 3.18287	<i>Elaeagnus glabra</i>	China	KX986779	KX999281	KX999251	KX999171	KX999212
<i>D. ellipicola</i>	CGMCC 3.17084	<i>Libocarpus glabra</i>	China	KF576270	N/A ^a	N/A ^a	KF576245	KF576291
<i>D. endophytica</i>	CBS 133811	<i>Schinus terebinthifolius</i>	Brazil	KC343065	KC343307	KC343549	KC343791	KC343065

Species	Isolate	Host	Location	GenBank accession numbers				
				ITS	cal	bis3	ref1	tub2
<i>D. eres</i>	AR5193	<i>Ulmus</i> sp.	Germany	KJ210529	KJ434999	KJ420850	KJ210550	KJ420799
	CFCC 52575	<i>Castanea mollissima</i>	China	MH121510	N/A ^a	MH121470	MH121552	MH121592
	CFCC 52576	<i>Castanea mollissima</i>	China	MH121511	MH121432	MH121471	MH121553	MH121593
	CFCC 52577	<i>Acanthopanax senticosus</i>	China	MH121512	MH121433	MH121472	MH121554	MH121594
	CFCC 52578	<i>Sorbus</i> sp.	China	MH121513	MH121434	MH121473	MH121555	MH121595
	CFCC 52579	<i>Juglans regia</i>	China	MH121514	N/A ^a	MH121474	MH121556	N/A ^a
	CFCC 52580	<i>Melia azedarace</i>	China	MH121515	N/A ^a	MH121475	MH121557	MH121596
	CFCC 52581	<i>Rhododendron simsii</i>	China	MH121516	N/A ^a	MH121476	MH121558	MH121597
	CBS 132525	<i>Eucalyptus</i> sp.	Australia	NR120157	N/Aa	N/Aa	N/Aa	N/Aa
<i>D. foeniculacea</i>	CBS 123208	<i>Foeniculum vulgare</i>	Portugal	KC343104	KC343346	KC343588	KC343830	KC344072
<i>D. fraxini-angustifoliae</i>	BRIP 54781	<i>Fraxinus angustifolia</i>	Australia	JX862528	N/A ^a	N/A ^a	JX862534	KF170920
<i>D. fraxinicola</i>	CFCC 52582	<i>Fraxinus chinensis</i>	China	MH121517	MH121435	N/A ^a	MH121559	N/A ^a
	CFCC 52583	<i>Fraxinus chinensis</i>	China	MH121518	MH121436	N/A ^a	MH121560	N/A ^a
<i>D. fukushii</i>	MAFF 625034	<i>Pyrus pyrifolia</i>	Japan	JQ807469	N/A ^a	N/A ^a	JQ807418	N/A ^a
<i>D. fusicola</i>	CGMCC 3.17087	<i>Lithocarpus glabra</i>	China	KF576281	KF576233	N/A ^a	KF576256	KF576305
<i>D. ganjae</i>	CBS 180.91	<i>Cannabis sativa</i>	USA	KC343112	KC343354	KC343596	KC343838	KC344080
<i>D. garebhjonesii</i>	MFLUCC 12-0542a	<i>Unknown dead leaf</i>	Thailand	KT459423	KT459470	N/A ^a	KT459457	KT459441
<i>D. goulteri</i>	BRIP 55657a	<i>Helianthus annuus</i>	Australia	KJ197290	N/A ^a	N/A ^a	KJ197252	KJ197270
<i>D. gulyae</i>	BRIP 54025	<i>Helianthus annuus</i>	Australia	JF431299	N/A ^a	N/A ^a	KJ197271	JN645803
<i>D. helianthi</i>	CBS 592.81	<i>Helianthus annuus</i>	Serbia	KC343115	KC343357	KC343599	KC343841	KC344083
<i>D. helicis</i>	AR5211	<i>Hedera helix</i>	France	KJ210538	KJ435043	KJ420875	KJ210559	KJ420828
<i>D. heterophyllae</i>	CBS 143769	<i>Acacia heterophylla</i>	France	MG600222	MG600218	MG600220	MG600224	MG600226
<i>D. hickoriae</i>	CBS 145.26	<i>Carya glabra</i>	USA	KC343118	KC343360	KC343602	KC343844	KC344086
<i>D. hispaniae</i>	CPC 30321	<i>Vitis vinifera</i>	Spain	MG281123	MG281820	MG281471	MG281644	MG281296
<i>D. hongkongensis</i>	CBS 115448	<i>Dichroa febrifuga</i>	China	KC343119	KC343361	KC343603	KC343845	KC344087
<i>D. incompleta</i>	CGMCC 3.18288	<i>Camellia sinensis</i>	China	KX986794	KX999289	KX999265	KX999186	KX999226
<i>D. inconspicua</i>	CBS 133813	<i>Maytenus ilicifolia</i>	Brazil	KC343123	KC343365	KC343607	KC343849	KC344091

Species	Isolate	Host	Location	GenBank accession numbers				
				ITS	cal	bis3	tefl	tub2
<i>D. infecunda</i>	CBS 133812	<i>Schinus terebinthifolius</i>	Brazil	KC343126	KC343368	KC343610	KC343852	KC344094
<i>D. isoberliniae</i>	CPC 22549	<i>Isoberlinia angolensis</i>	Zambia	KJ869133	N/A ^a	N/A ^a	N/A ^a	KJ869245
<i>D. juglandicola</i>	CFCC 51134	<i>Juglans mandshurica</i>	China	KU985101	KX024616	KX024622	KX024628	KX024634
	CFCC 51135	<i>Juglans mandshurica</i>	China	KU985102	KX024617	KX024623	KX024629	KX024635
<i>D. kadsurae</i>	CFCC 52586	<i>Kadsura longipedunculata</i>	China	MH121521	MH121439	MH121479	MH121563	MH121600
	CFCC 52587	<i>Kadsura longipedunculata</i>	China	MH121522	MH121440	MH121480	MH121564	MH121601
	CFCC 52588	<i>Acer</i> sp.	China	MH121523	MH121441	MH121481	MH121565	MH121602
	CFCC 52589	<i>Acer</i> sp.	China	MH121524	MH121442	MH121482	MH121566	MH121603
	BRIP 54033	<i>Helianthus annuus</i>	Australia	JF431295	N/A ^a	N/A ^a	JN645809	N/A ^a
<i>D. kochmanii</i>	BRIP 54031	<i>Portulaca grandiflora</i>	Australia	JF431301	N/A ^a	N/A ^a	JN645797	KJ197272
<i>D. litchicola</i>	BRIP 54900	<i>Litchi chinensis</i>	Australia	JX862533	N/A ^a	N/A ^a	JX862539	KF170925
<i>D. lithocarpus</i>	CGMCC 3.15175	<i>Lithocarpus glabra</i>	China	KC153104	KF576235	N/A ^a	KC153095	KF576311
<i>D. longicicola</i>	CGMCC 3.17089	<i>Lithocarpus glabra</i>	China	KF576267	N/A ^a	N/A ^a	KF576242	KF576291
<i>D. longicolla</i>	ATCC 60325	<i>Glycine max</i>	USA	KJ590728	N/A ^a	KJ659188	KJ590767	KJ610883
<i>D. longispora</i>	CBS 194.36	<i>Ribes</i> sp.	Canada	KC343135	KC343377	KC343619	KC343861	KC344103
<i>D. loniceriae</i>	MFLUCC 17-0963	<i>Lonicera</i> sp.	Italy	KY964190	KY964116	N/A ^a	KY964146	KY964073
<i>D. lusitanicae</i>	CBS 123212	<i>Foeniculum vulgare</i>	Portugal	KC343136	KC343378	KC343620	KC343862	KC344104
<i>D. macinthoshii</i>	BRIP 55064a	<i>Rapistrum rugostrum</i>	Australia	KJ197289	N/A ^a	N/A ^a	KJ197251	KJ197269
<i>D. mahothocarpus</i>	CGMCC 3.15181	<i>Lithocarpus glabra</i>	China	KC153096	N/A ^a	N/A ^a	KC153087	KF576312
<i>D. malorum</i>	CAA734	<i>Malus domestica</i>	Portugal	KY435638	KY435658	KY435648	KY435627	KY435668
<i>D. maritima</i>	DAOMC 250563	<i>Picea rubens</i>	Canada	N/A ^a	N/A ^a	N/A ^a	N/A ^a	KU574616
<i>D. masirevicii</i>	BRIP 57892a	<i>Helianthus annuus</i>	Australia	KJ197277	N/A ^a	N/A ^a	KJ197239	KJ197257
<i>D. mayteni</i>	CBS 133185	<i>Maytenus ilicifolia</i>	Brazil	KC343139	KC343381	KC343623	KC343865	KC344107
<i>D. maytenicola</i>	CPC 21896*	<i>Maytenus acuminata</i>	South Africa	KF777157	N/A ^a	N/A ^a	N/A ^a	KF777250
<i>D. melonis</i>	CBS 507.78	<i>Cucumis melo</i>	USA	KC343142	KC343384	KC343626	KC343868	KC344110
<i>D. middletonii</i>	BRIP 54884e	<i>Rapistrum rugostrum</i>	Australia	KJ197286	N/A ^a	N/A ^a	KJ197248	KJ197266
<i>D. miriciae</i>	BRIP 54736j	<i>Helianthus annuus</i>	Australia	KJ197282	N/A ^a	N/A ^a	KJ197244	KJ197262
<i>D. momicola</i>	MFLUCC 16-0113	<i>Prunus persica</i>	China	KU557563	KU557611	N/A ^a	KU557631	KU557558
<i>D. multiguttulata</i>	ZJUD98	<i>Citrus grandis</i>	China	KJ490633	N/A ^a	KJ490575	KJ490512	KJ490454

Species	Isolate	Host	Location	GenBank accession numbers				
				ITS	cal	bis3	ref1	tub2
<i>D. musigena</i>	CBS 129519	<i>Musa</i> sp.	Australia	KC343143	KC343385	KC343627	KC343869	KC344111
<i>D. neilliae</i>	CBS 144.27	<i>Spiraea</i> sp.	USA	KC343144	KC343386	KC343628	KC343870	KC344112
<i>D. neoarctii</i>	CBS 109490	<i>Ambrosia trifida</i>	USA	KC343145	KC343387	KC343629	KC343871	KC344113
<i>D. neoraonikayaporum</i>	MFLUCC 14-1136	<i>Tectona grandis</i>	Thailand	KU712449	KU749356	N/A ^a	KU749369	KU743988
<i>D. nobilis</i>	CBS 113470	<i>Castanea sativa</i>	Korea	KC343146	KC343388	KC343630	KC343872	KC344114
<i>D. nothofagi</i>	BRIP 54801	<i>Nothofagus cunninghamii</i>	Australia	JX862530	N/A ^a	N/A ^a	JX862536	KF170922
<i>D. novem</i>	CBS 127270	<i>Glycine max</i>	Croatia	KC343155	KC343397	KC343640	KC343881	KC344123
<i>D. ocoteae</i>	CBS 141330	<i>Ocotea obtusata</i>	France	KX228293	N/A ^a	N/A ^a	N/A ^a	KX228388
<i>D. oraccinii</i>	CGMCC 3.17531	<i>Camellia sinensis</i>	China	KP267863	N/A ^a	KP293517	KP267937	KP293443
<i>D. ovalispora</i>	ICMP20659	<i>Citrus limon</i>	China	KJ490628	N/A ^a	KJ490570	KJ490507	KJ490449
<i>D. ovoicicola</i>	CGMCC 3.17093	<i>Citrus</i> sp.	China	KF576265	KF576223	N/A ^a	KF576240	KF576289
<i>D. oxe</i>	CBS 133186	<i>Maytenus ilicifolia</i>	Brazil	KC343164	KC343406	KC343648	KC343890	KC344132
<i>D. padina</i>	CFCC 52590	<i>Padus racemosa</i>	China	MH121525	MH121443	MH121483	MH121567	MH121604
	CFCC 52591	<i>Padus racemosa</i>	China	MH121526	MH121444	MH121484	MH121568	MH121605
<i>D. pandanicola</i>	MFLU 18-0006	<i>Pandanus</i> sp.	Thailand	MG646974	N/A ^a	N/A ^a	N/A ^a	MG646930
<i>D. paranensis</i>	CBS 133184	<i>Maytenus ilicifolia</i>	Brazil	KC343171	KC343413	KC343655	KC343897	KC344139
<i>D. parapterocarpi</i>	CPC 22729	<i>Pterocarpus brenanii</i>	Zambia	KJ869138	N/A ^a	N/A ^a	N/A ^a	KJ869248
<i>D. pascoei</i>	BRIP 54847	<i>Persea americana</i>	Australia	JX862532	N/A ^a	N/A ^a	JX862538	KF170924
<i>D. passiflorae</i>	CBS 132527	<i>Passiflora edulis</i>	South America	JX069860	N/A ^a	KY435654	N/A ^a	N/A ^a
<i>D. passifloricola</i>	CBS 141329	<i>Passiflora foetida</i>	Malaysia	KX228292	N/A ^a	KX228367	N/A ^a	KX228387
<i>D. penetriteum</i>	CGMCC 3.17532	<i>Camellia sinensis</i>	China	KP714505	N/A ^a	KP714493	KP714517	KP714529
<i>D. perijuncta</i>	CBS 109745	<i>Ulmus glabra</i>	Austria	KC343172	KC343414	KC343656	KC343898	KC344140
<i>D. perseae</i>	CBS 151.73	<i>Persea gratissima</i>	Netherlands	KC343173	KC343415	KC343657	KC343899	KC344141
<i>D. pescicola</i>	MFLUCC 16-0105	<i>Prunus persica</i>	China	KU557555	KU557603	N/A ^a	KU557623	KU557579
<i>D. phaseolorum</i>	AR4203	<i>Phaseolus vulgaris</i>	USA	KJ590738	N/A ^a	KJ659220	N/A ^a	KP004507
<i>D. podocarpi-macrophylli</i>	CGMCC 3.18281	<i>Podocarpus macrophyllus</i>	China	KX986774	KX999278	KX999246	KX999167	KX999207
<i>D. pseudomangiferae</i>	CBS 101339	<i>Mangifera indica</i>	Dominican Republic	KC343181	KC343423	KC343665	KC343907	KC344149

Species	Isolate	Host	Location	GenBank accession numbers				
				ITS	cal	his3	tefl	tub2
<i>D. pseudophoenicicola</i>	CBS 462.69	<i>Phoenix dactylifera</i>	Spain	KC343184	KC343426	KC343668	KC343910	KC344152
<i>D. pseudotsugae</i>	MFLU 15-3228	<i>Pseudotsuga menziesii</i>	Italy	KY964225	KY964138	N/A ^a	KY964181	KY964108
<i>D. psoraleae</i>	CBS 136412	<i>Psoralea pinnata</i>	South Africa	KF777158	N/A ^a	N/A ^a	KF777245	KF777251
<i>D. psoraleae-pinnatae</i>	CBS 136413	<i>Psoralea pinnata</i>	South Africa	KF777159	N/A ^a	N/A ^a	N/A ^a	KF777252
<i>D. pterocarpi</i>	MFLUCC 10-0571	<i>Pterocarpus indicus</i>	Thailand	JQ619899	JX197451	N/A ^a	JX275416	JX275460
<i>D. pterocarpicola</i>	MFLUCC 10-0580a	<i>Pterocarpus indicus</i>	Thailand	JQ619887	JX197433	N/A ^a	JX275403	JX275441
<i>D. pulla</i>	CBS 338.89	<i>Hedera helix</i>	Yugoslavia	KC343152	KC343394	KC343636	KC343878	KC344120
<i>D. pyracanthae</i>	CAA483	<i>Pyracantha coccinea</i>	Portugal	KY435635	KY435656	KY435645	KY435625	KY435666
<i>D. racemosae</i>	CBS 143770	<i>Euclea racemosa</i>	South Africa	MG600223	MG600219	MG600221	MG600225	MG600227
<i>D. raonikayaporum</i>	CBS 133182	<i>Spondias mombin</i>	Brazil	KC343188	KC343430	KC343672	KC343914	KC344156
<i>D. ravennica</i>	MFLUCC 15-0479	<i>Tamarix</i> sp.	Italy	KU900335	N/A ^a	N/A ^a	KX365197	KX432254
<i>D. rhusicola</i>	CBS 129528	<i>Rhus pendulina</i>	South Africa	JF951146	KC843124	N/A ^a	KC843100	KC843205
<i>D. rosae</i>	MFLU 17-1550	<i>Rosa</i> sp.	Thailand	MG828894	N/A ^a	N/A ^a	N/A ^a	MG843878
<i>D. rosicola</i>	MFLU 17-0646	<i>Rosa</i> sp.	UK	MG828895	N/A ^a	N/A ^a	MG829270	MG843877
<i>D. rostrata</i>	CFCC 50062	<i>Juglans mandshurica</i>	China	KP208847	KP208849	KP208851	KP208853	KP208855
	CFCC 50063	<i>Juglans mandshurica</i>	China	KP208848	KP208850	KP208852	KP208854	KP208856
<i>D. rudis</i>	AR3422	<i>Laburnum anagyroides</i>	Austria	KC843331	KC843146	N/A ^a	KC843090	KC843177
<i>D. saccharata</i>	CBS 116311	<i>Protea repens</i>	South Africa	KC343190	KC343432	KC343674	KC343916	KC344158
<i>D. sackstonii</i>	BRIP 54669b	<i>Helianthus annuus</i>	Australia	KJ197287	N/A ^a	N/A ^a	KJ197249	KJ197267
<i>D. salicicola</i>	BRIP 54825	<i>Salix purpurea</i>	Australia	JX862531	N/A ^a	N/A ^a	JX862537	JX862531
<i>D. sambucusii</i>	CFCC 51986	<i>Sambucus williamsii</i>	China	KY852495	KY852499	KY852503	KY852507	KY852511
	CFCC 51987	<i>Sambucus williamsii</i>	China	KY852496	KY852500	KY852504	KY852508	KY852512
<i>D. schini</i>	CBS 133181	<i>Schinus terebinthifolius</i>	Brazil	KC343191	KC343433	KC343675	KC343917	KC344159
<i>D. schisandrae</i>	CFCC 51988	<i>Schisandra chinensis</i>	China	KY852497	KY852501	KY852505	KY852509	KY852513
	CFCC 51989	<i>Schisandra chinensis</i>	China	KY852498	KY852502	KY852506	KY852510	KY852514
<i>D. schoeni</i>	MFLU 15-1279	<i>Schoenus nigricans</i>	Italy	KY964226	KY964139	N/A ^a	KY964182	KY964109
<i>D. sclerotioides</i>	CBS 296.67	<i>Cucumis sativus</i>	Netherlands	KC343193	KC343435	KC343677	KC343919	KC344161

Species	Isolate	Host	Location	GenBank accession numbers				
				ITS	cal	bis3	ref1	tub2
<i>D. sennae</i>	CFCC 51636	<i>Senna bicapsularis</i>	China	KY203724	KY228875	N/A ^a	KY228885	KY228891
	CFCC 51637	<i>Senna bicapsularis</i>	China	KY203725	KY228876	N/Aa	KY228886	KY228892
<i>D. sennicola</i>	CFCC 51634	<i>Senna bicapsularis</i>	China	KY203722	KY228873	KY228879	KY228883	KY228889
	CFCC 51635	<i>Senna bicapsularis</i>	China	KY203723	KY228874	KY228880	KY228884	KY228890
<i>D. serafiniae</i>	BRIP 55665a	<i>Helianthus annuus</i>	Australia	KJ197274	N/A ^a	N/A ^a	KJ197236	KJ197254
<i>D. siamensis</i>	MFLUCC 10-573a	<i>Dasymaschalon</i> sp.	Thailand	JQ619879	N/A ^a	N/A ^a	JX275393	JX275429
<i>D. sojae</i>	FAU635	<i>Glycine max</i>	USA	KJ590719	KJ612116	KJ659208	KJ590762	KJ610875
<i>D. spartinicola</i>	CBS 140003	<i>Spartium junceum</i>	Spain	KR611879	N/A ^a	KR857696	N/A ^a	KR857695
<i>D. sterilis</i>	CBS 136969	<i>Vaccinium corymbosum</i>	Italy	KJ160579	KJ160548	MF418350	KJ160611	KJ160528
<i>D. stictica</i>	CBS 370.54	<i>Buxus sampervirens</i>	Italy	KC343212	KC343454	KC343696	KC343938	KC344180
<i>D. subclavata</i>	ICMP20663	<i>Citrus unshiu</i>	China	KJ490587	N/A ^a	KJ490529	KJ490466	KJ490408
<i>D. subcylindrospora</i>	MFLU 17-1195	<i>Salix</i> sp.	China	MG746629	N/A ^a	N/A ^a	MG746630	MG746631
<i>D. subellipicola</i>	MFLU 17-1197	on dead wood	China	MG746632	N/A ^a	N/A ^a	MG746633	MG746634
<i>D. subordinaria</i>	CBS 464.90	<i>Plantago lanceolata</i>	New Zealand	KC343214	KC343456	KC343698	KC343940	KC344182
<i>D. taoicola</i>	MFLUCC 16-0117	<i>Prunus persica</i>	China	KU557567	N/Aa	N/A ^a	KU557635	KU557591
<i>D. tectonae</i>	MFLUCC 12-0777	<i>Tectona grandis</i>	China	KU712430	KU749345	N/A ^a	KU749359	KU743977
<i>D. tectonendophytica</i>	MFLUCC 13-0471	<i>Tectona grandis</i>	China	KU712439	KU749354	N/A ^a	KU749367	KU749354
<i>D. tectonigena</i>	MFLUCC 12-0767	<i>Tectona grandis</i>	China	KU712429	KU749358	N/A ^a	KU749371	KU743976
<i>D. terebinthifolii</i>	CBS 133180	<i>Schinus terebinthifolius</i>	Brazil	KC343216	KC343458	KC343700	KC343942	KC344184
<i>D. thunbergii</i>	MFLUCC 10-576a	<i>Thunbergia laurifolia</i>	Thailand	JQ619893	JX197440	N/A ^a	JX275409	JX275449
<i>D. thunbergiicola</i>	MFLUCC 12-0033	<i>Thunbergia laurifolia</i>	Thailand	KP715097	N/A ^a	N/A ^a	KP715098	N/A ^a
<i>D. tibetensis</i>	CFCC 51999	<i>Juglandis regia</i>	China	MF279843	MF279888	MF279828	MF279858	MF279873
	CFCC 52000	<i>Juglandis regia</i>	China	MF279844	MF279889	MF279829	MF279859	MF279874
<i>D. torilicola</i>	MFLUCC 17-1051	<i>Torilis arvensis</i>	Italy	KY964212	KY964127	N/A ^a	KY964168	KY964096
<i>D. toxica</i>	CBS 534.93	<i>Lupinus angustifolius</i>	Australia	KC343220	KC343462	C343704	KC343946	KC344188
<i>D. tulliensis</i>	BRIP 62248a	<i>Theobroma cacao</i> fruit	Australia	KR936130	N/A ^a	N/A ^a	KR936133	KR936132
<i>D. ueckerae</i>	FAU656	<i>Cucumis melo</i>	USA	KJ590726	KJ612122	KJ659215	KJ590747	KJ610881
<i>D. ukurunduensis</i>	CFCC 52592	<i>Acer ukurunduense</i>	China	MH121527	MH121445	MH121485	MH121569	N/A ^a
	CFCC 52593	<i>Acer ukurunduense</i>	China	MH121528	MH121446	MH121486	MH121570	N/A ^a

Species	Isolate	Host	Location	GenBank accession numbers				
				ITS	cal	bis3	tef1	tub2
<i>D. undulata</i>	CGMCC 3.18293	Leaf of unknown host	China-Laos border	KX986798	N/A ^a	KX999269	KX999190	KX999230
	CGMCC 3.17569	<i>Citrus unshiu</i>	China	KJ490587	N/A ^a	KJ490529	KJ490408	KJ490466
<i>D. unshiuensis</i>	CFCC 52594	<i>Carya illinoensis</i>	China	MH121529	MH121447	MH121487	MH121571	MH121606
	CFCC 52595	<i>Carya illinoensis</i>	China	MH121530	MH121448	MH121488	MH121572	MH121607
<i>D. vaccinii</i>	CBS 160.32	<i>Oxycoccus macrocarpos</i>	USA	KC343228	KC343470	KC343712	KC343954	KC344196
<i>D. vanguardiae</i>	CPC 22703	<i>Vangueria infausta</i>	Zambia	KJ869137	N/A ^a	N/A ^a	N/A ^a	KJ869247
<i>D. vawdreyi</i>	BRIP 57887a	<i>Psidium guajava</i>	Australia	KR936126	N/A ^a	N/A ^a	KR936129	KR936128
<i>D. velutina</i>	CGMCC 3.18286	<i>Neolitsea</i> sp.	China	KX986790	N/A ^a	KX999261	KX999182	KX999223
<i>D. virgiliae</i>	CMW/40748	<i>Virgilia oroboides</i>	South Africa	KP247566	N/A ^a	N/A ^a	N/A ^a	KP247575
<i>D. xishuangbanica</i>	CGMCC 3.18282	<i>Camellia sinensis</i>	China	KX986783	N/A ^a	KX999255	KX999175	KX999216
<i>D. yunnanensis</i>	CGMCC 3.18289	<i>Coffea</i> sp.	China	KX986796	KX999290	KX999267	KX999188	KX999228
<i>Diaporthella corylina</i>	CBS 121124	<i>Corylus</i> sp.	China	KC343004	KC343246	KC343488	KC343730	KC343972

Newly sequenced material is indicated in bold type.

Table 2. Genes used in this study with PCR primers, process and references.

Gene	PCR primers (forward/reverse)	PCR: thermal cycles: (Annealing temp. in bold)	References of primers used
ITS	ITS1/ITS4	(95 °C: 30 s, 51 °C : 30 s, 72 °C: 1 min) × 35 cycles	White et al. 1990
cal	CAL228F/CAL737R	(95 °C: 15 s, 55 °C : 20 s, 72 °C: 1 min) × 35 cycles	Carbone and Kohn 1999
his3	CYLH4F/H3-1b	(95 °C: 30 s, 58 °C : 30 s, 72 °C: 1 min) × 35 cycles	Glass and Donaldson 1995, Crous et al. 2004a
tef1	EF1-728F/EF1-986R	(9 °C: 15 s, 55 °C : 20 s, 72 °C: 1 min) × 35 cycles	Carbone and Kohn 1999
tub2	T1(Bt2a)/Bt2b	(95 °C: 30 s, 55 °C : 30 s, 72 °C: 1 min) × 35 cycles	Glass and Donaldson 1995, Glass and Donaldson 1995

Materials and methods

Isolates

From 2015 to 2017, fresh specimens of *Diaporthe* were collected from symptomatic or non-symptomatic twigs or branches from Beijing, Heilongjiang, Jiangsu, Jiangxi, Shaanxi and Zhejiang Provinces in China (Table 1). A total of 105 isolates were obtained by removing a mucoid spore mass from conidiomata and spreading the suspension on the surface of 1.8% potato dextrose agar (PDA) in a Petri dish and incubating at 25 °C for up to 24 h. Single germinating conidia were transferred on to fresh PDA plates. Forty-two representative *Diaporthe* strains were selected based on cultural characteristics on PDA, conidia morphology and ITS sequence data. Specimens were deposited in the Museum of the Beijing Forestry University (BJFC). Axenic cultures are maintained in the China Forestry Culture Collection Centre (CFCC).

Morphological analysis

Agar plugs (6 mm diam.) were taken from the edge of actively growing cultures on PDA and transferred on to the centre of 9 cm diam Petri dishes containing 2% tap water agar supplemented with sterile pine needles (PNA; Smith et al. 1996) and potato dextrose agar (PDA) and incubated at 20–21 °C under a 12 h near-ultraviolet light/12 h dark cycle to induce sporulation as described in recent studies (Gomes et al. 2013, Lombard et al. 2014). Colony characters and pigment production on PNA and PDA were noted after 10 d. Colony colours were rated according to Rayner (1970). Cultures were examined periodically for the development of ascomata and conidiomata. The morphological characteristics were examined by mounting fungal structures in clear lactic acid and 30 measurements at 1000× magnification were determined for each isolate using a Leica compound microscope (DM 2500) with interference contrast (DIC) optics. Descriptions, nomenclature and illustrations of taxonomic novelties are deposited in MycoBank (www.MycoBank.org; Crous et al. 2004b).

DNA extraction, PCR amplification and sequencing

Genomic DNA was extracted from colonies grown on cellophane-covered PDA using a modified CTAB [cetyltrimethylammonium bromide] method (Doyle and Doyle 1990). DNA was estimated by electrophoresis in 1% agarose gel and the quality was measured using the NanoDrop 2000 (Thermo Scientific, Waltham, MA, USA), following the user manual (Desjardins et al. 2009). PCR amplifications were performed in a DNA Engine Peltier Thermal Cycler (PTC-200; Bio-Rad Laboratories, Hercules, CA, USA). The primer sets ITS1/ITS4 (White et al. 1990) were used to amplify the ITS region. The primer pair CAL228F/CAL737R (Carbone and Kohn 1999) were

used to amplify the calmodulin gene (*cal*) and the primer pair CYLH4F (Crous et al. 2004a) and H3-1b (Glass and Donaldson 1995) were used to amplify part of the histone H3 (*his3*) gene. The primer pair EF1-728F/EF1-986R (Carbone and Kohn 1999) were used to amplify a partial fragment of the translation elongation factor 1- α gene (*tef1*). The primer sets T1 (O'Donnell and Cigelnik 1997) and Bt2b (Glass and Donaldson 1995) were used to amplify the beta-tubulin gene (*tub2*); the additional combination of Bt2a/Bt2b (Glass and Donaldson 1995) was used in case of amplification failure of the T1/Bt2b primer pair. Amplifications of different loci were performed under different conditions (Table 2). PCR amplification products were assayed via electrophoresis in 2% agarose gels. DNA sequencing was performed using an ABI PRISM® 3730XL DNA Analyser with a BigDye Terminator Kit v.3.1 (Invitrogen, USA) at the Shanghai Invitrogen Biological Technology Company Limited (Beijing, China).

Phylogenetic analyses

DNA generated sequences were used to obtain consensus sequences using SeqMan v.7.1.0 DNASTAR Lasergene Core Suite software programme (DNASTAR Inc., Madison, WI, USA). Sequences were aligned using MAFFT v.6 (Kato and Toh 2010) and edited manually using MEGA6 (Tamura et al. 2013). Two different datasets were employed to estimate two phylogenetic analyses: one for *Diaporthe* species and one for *Diaporthe eres* complex. The first analysis was undertaken to infer the interspecific relationships in *Diaporthe*. All the *Diaporthe* isolates recovered from samples collected during this study and additional reference sequences of *Diaporthe* species were included in the dataset of combined ITS, *cal*, *his3*, *tef1*, and *tub2* regions (Table 1), with *Diaporthella corylina* (CBS 121124) as outgroup. The second analysis focused on the *Diaporthe eres* complex based on *cal*, *tef1* and *tub2* loci (Table 3) according to recent publications (Gao et al. 2014, 2015, 2016, Udayanga et al. 2014b, Tanney et al. 2016, Fan et al. 2018), with *Diaporthe citri* (AR3405) as outgroup. Maximum Parsimony analysis was performed by a heuristic search option of 1000 random-addition sequences with a tree bisection and reconnection (TBR) algorithm. Maxtrees were set to 5000, branches of zero length were collapsed and all equally parsimonious trees were saved. Other calculated parsimony scores were tree length (TL), consistency index (CI), retention index (RI) and rescaled consistency (RC). Maximum Likelihood analysis was performed with a GTR site substitution model (Guindon et al. 2010). Branch support was evaluated with a bootstrapping (BS) method of 1000 replicates (Hillis and Bull 1993).

Bayesian inference (BI) analysis, employing a Markov chain Monte Carlo (MCMC) algorithm, was performed (Rannala and Yang 1996). MrModeltest v. 2.3 was used to estimate the best-fit model of nucleotide substitution model settings for each gene (Posada and Crandall 1998). Two MCMC chains started from random trees for 1,000,000 generations and trees were sampled every 100th generation, resulting in

Table 3. Isolates and GenBank accession numbers used in the phylogenetic analyses of *Diaporthe eres* complex.

Species	Isolate/culture collection	Host	Location	GenBank accession numbers		
				CAL	TEF1- α	TUB
<i>D. alleghaniensis</i>	CBS 495.72	<i>Betula alleghaniensis</i>	Canada	KC343249	GQ250298	KC843228
<i>D. alnea</i>	CBS 146.46	<i>Alnus</i> sp.	Netherlands	KC343250	KC343734	KC343976
	CBS 159.47	<i>Alnus</i> sp.	Netherlands	KC343251	KC343735	KC343977
	LCM22b.02a	<i>Alnus</i> sp.	USA	KJ435020	KJ210557	KJ420825
	LCM22b.02b	<i>Alnus</i> sp.	USA	KJ435021	KJ210558	KJ420826
<i>D. betulina</i>	CFCC 52560	<i>Betula albo-sinensis</i>	China	MH121419	MH121537	MH121577
	CFCC 52561	<i>Betula costata</i>	China	MH121420	MH121538	MH121578
	CFCC 52562	<i>Betula platyphylla</i>	China	MH121421	MH121539	MH121579
<i>D. bicincta</i>	CBS 121004	<i>Juglans</i> sp.	USA	KC343376	KC343860	KC344102
<i>D. biguttusis</i>	CGMCC 3.17081	<i>Lithocarpus glabra</i>	China	N/A ^a	KF576257	KF576306
<i>D. camptothecicola</i>	CFCC 51632	<i>Camptotheca acuminata</i>	China	KY228881	KY228887	KY228893
<i>D. celastrina</i>	CBS 139.27	<i>Celastrus</i> sp.	USA	KC343289	KC343773	KC344015
<i>D. chensiensis</i>	CFCC 52567	<i>Abies chensiensis</i>	China	MH121426	MH121544	MH121584
	CFCC 52568	<i>Abies chensiensis</i>	China	MH121427	MH121545	MH121585
<i>D. citri</i>	AR3405	<i>Citrus</i> sp.	USA	KC843157	KC843071	KC843187
<i>D. citrichinensis</i>	ZJUD034	<i>Citrus</i> sp.	China	KC843234	KC843071	KC843187
	ZJUD034B	<i>Citrus</i> sp.	China	KJ435042	KJ210562	KJ420829
<i>D. ellipicola</i>	CGMCC 3.17084	<i>Lithocarpus glabra</i>	China	N/A ^a	KF576245	KF576291
<i>D. eres</i>	AR5193	<i>Ulmus laevis</i>	Germany	KJ434999	KJ210550	KJ420799
	AR5196	<i>Ulmus laevis</i>	Germany	KJ435006	KJ210554	KJ420817
	DP0438	<i>Ulmus minor</i>	Austria	KJ435016	KJ210553	KJ420816
	LCM114.01a	<i>Ulmus</i> sp.	USA	KJ435027	KJ210545	KJ420787
	LCM114.01b	<i>Ulmus</i> sp.	USA	KJ435026	KJ210544	KJ420786
	FAU483	<i>Malus</i> sp.	Netherlands	KJ435022	JQ807422	KJ420827
	DAN001A	<i>Daphne laureola</i>	France	KJ434994	KJ210540	KJ420781
	DAN001B	<i>Daphne laureola</i>	France	KJ434995	KJ210541	KJ420782
	AR5197	<i>Rhododendron</i> sp.	Germany	KJ435014	KJ210552	KJ420812
	CBS 439.82	<i>Cotoneaster</i> sp.	UK	JX197429	GQ250341	JX275437
	AR3519	<i>Corylus avellana</i>	Austria	KJ435008	KJ210547	KJ420789
	FAU506	<i>Cornus florida</i>	USA	KJ435012	JQ807403	KJ420792
	FAU570	<i>Oxydendrum arboreum</i>	USA	KJ435025	JQ807410	KJ420794
	AR3723	<i>Rubus fruticosus</i>	Austria	KJ435024	JQ807354	KJ420793
	FAU522	<i>Sassafras albida</i>	USA	KJ435010	JQ807406	KJ420791
	DP0666	<i>Juglans cinerea</i>	USA	KJ435007	KJ210546	KJ420788
	DP0667	<i>Juglans cinerea</i>	USA	KC843155	KC843121	KC843229
	AR3560	<i>Viburnum</i> sp.	Austria	KJ435011	JQ807351	KJ420795
	AR5224	<i>Hedera helix</i>	Germany	KJ435036	KJ210551	KJ420802
	AR5231	<i>Hedera helix</i>	Germany	KJ435038	KJ210555	KJ420818
	AR5223	<i>Acer nugundo</i>	Germany	KJ435000	KJ210549	KJ420830
	CBS 109767	<i>Acer</i> sp.	Austria	KC343317	KC343801	KC344043
	DLR12a	<i>Vitis vinifera</i>	France	KJ434996	KJ210542	KJ420783
	DLR12b	<i>Vitis vinifera</i>	France	KJ434997	KJ210543	KJ420784
	AR4347	<i>Vitis vinifera</i>	Korea	KJ435030	JQ807356	KJ420805
	AR4355	<i>Prunus</i> sp.	Korea	KJ435035	JQ807359	KJ420797
	AR4367	<i>Prunus</i> sp.	Korea	KJ435019	JQ807364	KJ420824
	AR4346	<i>Prunus mume</i>	Korea	KJ435003	JQ807355	KJ420823
	AR4348	<i>Prunus persici</i>	Korea	KJ435004	JQ807357	JQ807357
	AR3669	<i>Pyrus pyrifolia</i>	Japan	KJ435002	JQ807415	KJ420808
	AR3670	<i>Pyrus pyrifolia</i>	Japan	KJ435001	JQ807416	KJ420807
	AR3671	<i>Pyrus pyrifolia</i>	Japan	KJ435017	JQ807417	KJ420814

Species	Isolate/culture collection	Host	Location	GenBank accession numbers		
				CAL	TEF1- α	TUB
	AR3672	<i>Pyrus pyrifolia</i>	Japan	KJ435023	JQ807418	KJ420819
	DP0591	<i>Pyrus pyrifolia</i>	New Zealand	KJ435018	JQ807395	KJ420821
	AR4369	<i>Pyrus pyrifolia</i>	Korea	KJ435005	JQ807366	KJ420813
	DP0180	<i>Pyrus pyrifolia</i>	New Zealand	KJ435029	JQ807384	KJ420804
	DP0179	<i>Pyrus pyrifolia</i>	New Zealand	KJ435028	JQ807383	KJ420803
	DP0590	<i>Pyrus pyrifolia</i>	New Zealand	KJ435037	JQ807394	KJ420810
	AR4373	<i>Ziziphus jujuba</i>	Korea	KJ435013	JQ807368	KJ420798
	AR4374	<i>Ziziphus jujuba</i>	Korea	KJ434998	JQ807369	KJ420785
	AR4357	<i>Ziziphus jujuba</i>	Korea	KJ435031	JQ807360	KJ420806
	AR4371	<i>Malus pumila</i>	Korea	KJ435034	JQ807367	KJ420796
	FAU532	<i>Chamaecyparis thyoides</i>	USA	KJ435015	JQ807408	KJ435015
	CBS 113470	<i>Castanea sativa</i>	Australia	KC343388	KC343872	KC344114
	AR4349	<i>Vitis vinifera</i>	Korea	KJ435032	JQ807358	KJ420822
	AR4363	<i>Malus</i> sp.	Korea	KJ435033	JQ807362	KJ420809
	CFCC 52575	<i>Castanea mollissima</i>	China	N/A^a	MH121552	MH121592
	CFCC 52576	<i>Castanea mollissima</i>	China	MH121432	MH121553	MH121593
	CFCC 52577	<i>Acanthopanax senticosus</i>	China	MH121433	MH121554	MH121594
	CFCC 52578	<i>Sorbus</i> sp.	China	MH121434	MH121555	MH121595
<i>D. helici</i>	CFCC 52579	<i>Juglans regia</i>	China	N/A^a	MH121556	N/A^a
	CFCC 52580	<i>Melia azedarace</i>	China	N/A^a	MH121557	MH121596
	CFCC 52581	<i>Rhododendron simsii</i>	China	N/A^a	MH121558	MH121597
	AR5211	<i>Hedera helix</i>	France	KJ435043	KJ210559	KJ420828
	CGMCC 3.17089	<i>Lithocarpus glabra</i>	China	N/A ^a	KF576242	KF576291
	CGMCC 3.15181	<i>Lithocarpus glabra</i>	China	N/A ^a	KC153087	KF576312
	DAOMC 250563	<i>Picea rubens</i>	Canada	N/A ^a	N/A ^a	KU574616
	MFLUCC 16-0113	<i>Prunus persica</i>	China	N/A ^a	KU557631	KU55758
	CBS 144. 27	<i>Spiraea</i> sp.	USA	KC343386	KC343870	KC344112
	CFCC 52590	<i>Padus racemosa</i>	China	MH121443	MH121567	MH121604
	CFCC 52591	<i>Padus racemosa</i>	China	MH121444	MH121568	MH121605
	CBS 138897	<i>Phragmites australis</i>	China	N/A ^a	N/A ^a	KP004507
	CBS 338.89	<i>Hedera helix</i>	Yugoslavia	KC343394	KC343878	KC344120
<i>D. padina</i>	DF5032	<i>Vaccinium corymbosum</i>	USA	KC849457	JQ807380	KC843225
	FAU633	<i>Vaccinium macrocarpon</i>	USA	KC849456	JQ807413	KC843226
	FAU446	<i>Vaccinium macrocarpon</i>	USA	KC849455	JQ807398	KC843224
	CBS 160.32	<i>Vaccinium macrocarpon</i>	USA	KC343470	GQ250326	JX270436
	FAU 468	<i>Vaccinium macrocarpon</i>	USA	KC849458	JQ807399	KC843227

Newly sequenced material is indicated in bold type.

a total of 10,000 trees. The first 25% of trees were discarded as the burn-in phase of each analysis. Branches with significant Bayesian posterior probabilities (BPP) were estimated in the remaining 7500 trees.

Sequences data were deposited in GenBank (Table 1). The multilocus sequence alignments were deposited in TreeBASE (www.treebase.org) as accession S22702 and S22703. The taxonomic novelties were deposited in MycoBank (Crous et al. 2004b).

Results

Collection of *Diaporthe* strains

Forty-two representative *Diaporthe* strains were isolated from 16 different host genera (Table 1) collected from six Provinces (Beijing, Heilongjiang, Jiangsu, Jiangxi, Shaanxi and Zhejiang) in China. All of these strains were isolated from symptomatic or non-symptomatic branches or twigs and preserved in the China Forestry Culture Collection Centre (CFCC).

Phylogenetic analyses

The first sequences dataset for the ITS, *cal*, *his3*, *tef1*, and *tub2* was analysed in combination to infer the interspecific relationships within *Diaporthe*. The combined species phylogeny of the *Diaporthe* isolates consisted of 236 sequences, including the outgroup sequences of *Diaporthella corylina* (culture CBS 121124). A total of 2948 characters including gaps (516 for ITS, 568 for *cal*, 520 for *his3*, 486 for *tef1* and 858 for *tub2*) were included in the phylogenetic analysis. The maximum likelihood tree, conducted by the GTR model, confirmed the tree topology and posterior probabilities of the Bayesian consensus tree. For the Bayesian analyses, MrModeltest suggested that all partitions should be analysed with dirichlet state frequency distributions. The following models were recommended by MrModeltest and used: GTR+I+G for ITS, *cal* and *his3*, HKY+I+G for *tef1* and *tub2*. The topology and branching order of ML were similar to BI analyses (Fig. 1). Based on the multi-locus phylogeny and morphology, 42 strains were assigned to 15 species, including 12 taxa which we describe here as new (Fig. 1).

The second dataset with *cal*, *tef1* and *tub2* sequences were analysed to focus on the *Diaporthe eres* complex. The alignment included 86 taxa, including the outgroup sequences of *Diaporthe citri* (Table 3). The aligned three-locus datasets included 1148 characters. Of these, 881 characters were constant, 105 variable characters were parsimony-uninformative and 162 characters were parsimony informative. The heuristic search using maximum parsimony (MP) generated 105 parsimonious trees (TL = 438, CI = 0.669, RI = 0.883, RC = 0.591), from which one was selected (Fig. 2). Based on the multi-locus phylogeny and morphology, seven strains were identified as *D. eres*, seven strains formed three distinct clades embedded in the *D. eres* complex, i.e. *D. betulina*, *D. chensiensis* and *D. padina*. MP and ML bootstrap support values above 50% are shown as first and second position, respectively. The branches with significant Bayesian posterior probability (≥ 0.70) in Bayesian analyses were thickened in the phylogenetic tree. The current results, based on the three genes (*cal*, *tef1* and *tub2*), suggest that *D. eres* clade could be separated from other species in this complex (Fig. 2). However, *D. biguttusis* (CGMCC 3.17081), *D. camptothecicola* (CFCC 51632), *D. ellipicola* (CGMCC 3.17084), *D. longicicola* (CGMCC 3.17089), *D. mahothocarpus* (CGMCC 3.15181) and *D. momicola* (MFLUCC 16-0113) were clustered in *D. eres* clade and thus treated as the synonyms of *D. eres* in the current study.

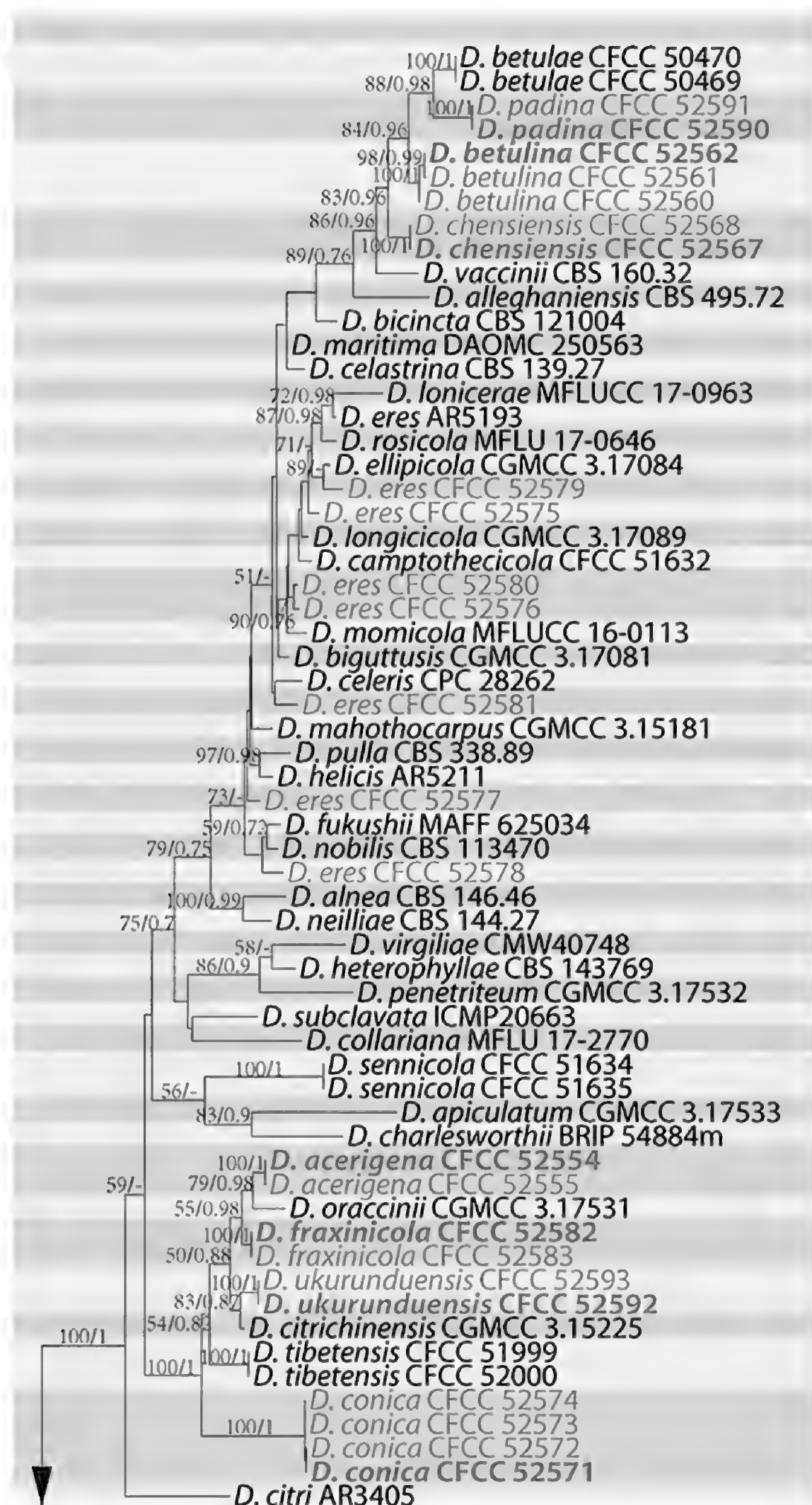


Figure 1. Phylogram of *Diaporthe* from a maximum likelihood analysis based on combined ITS, *cal*, *his3*, *tef1* and *tub2*. Values above the branches indicate maximum likelihood bootstrap (left, ML BP \geq 50%) and bayesian probabilities (right, BI PP \geq 0.70). The tree is rooted with *Diaporthella corylina*. Strains in the current study are in blue.

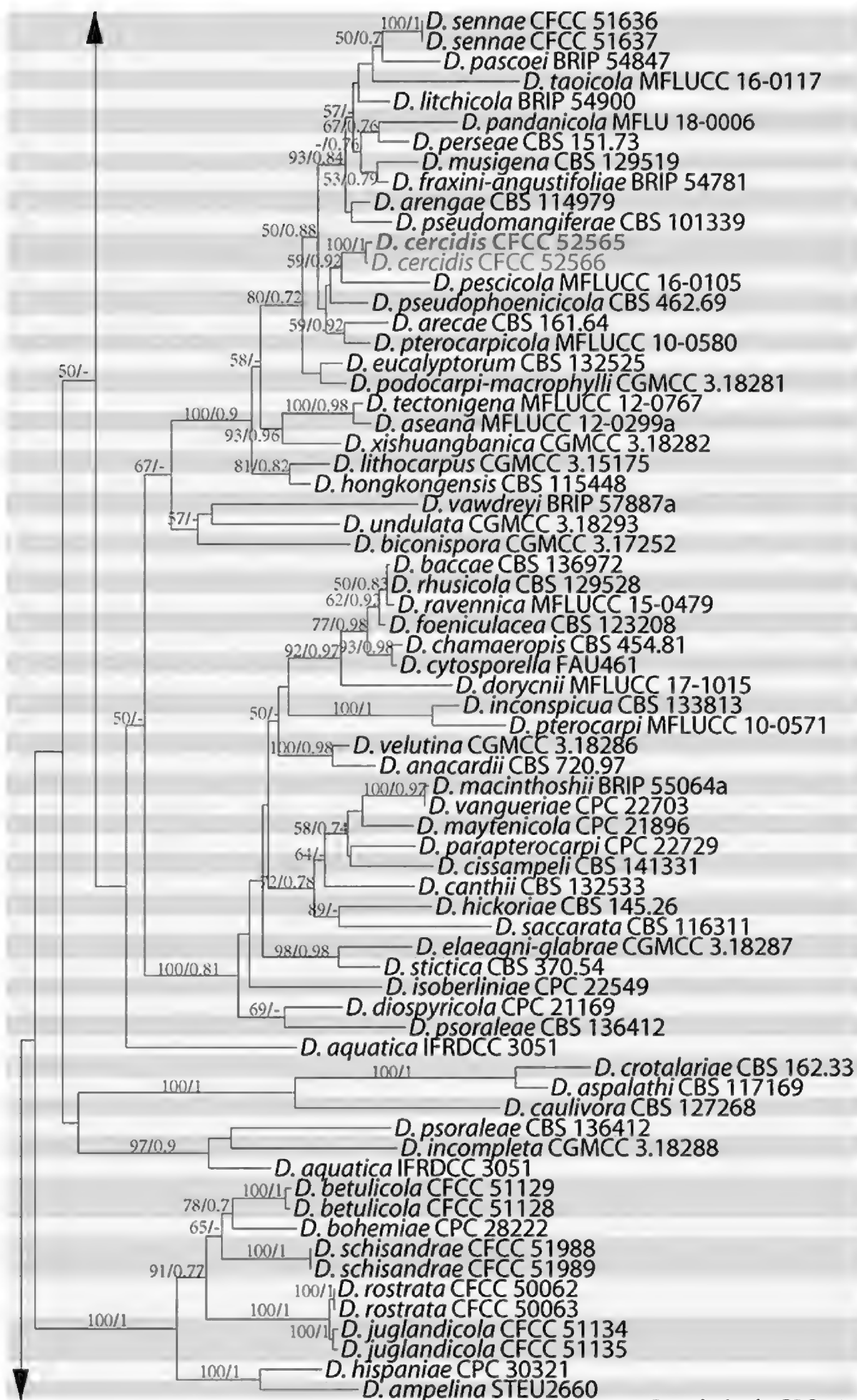


Figure 1. Continued.

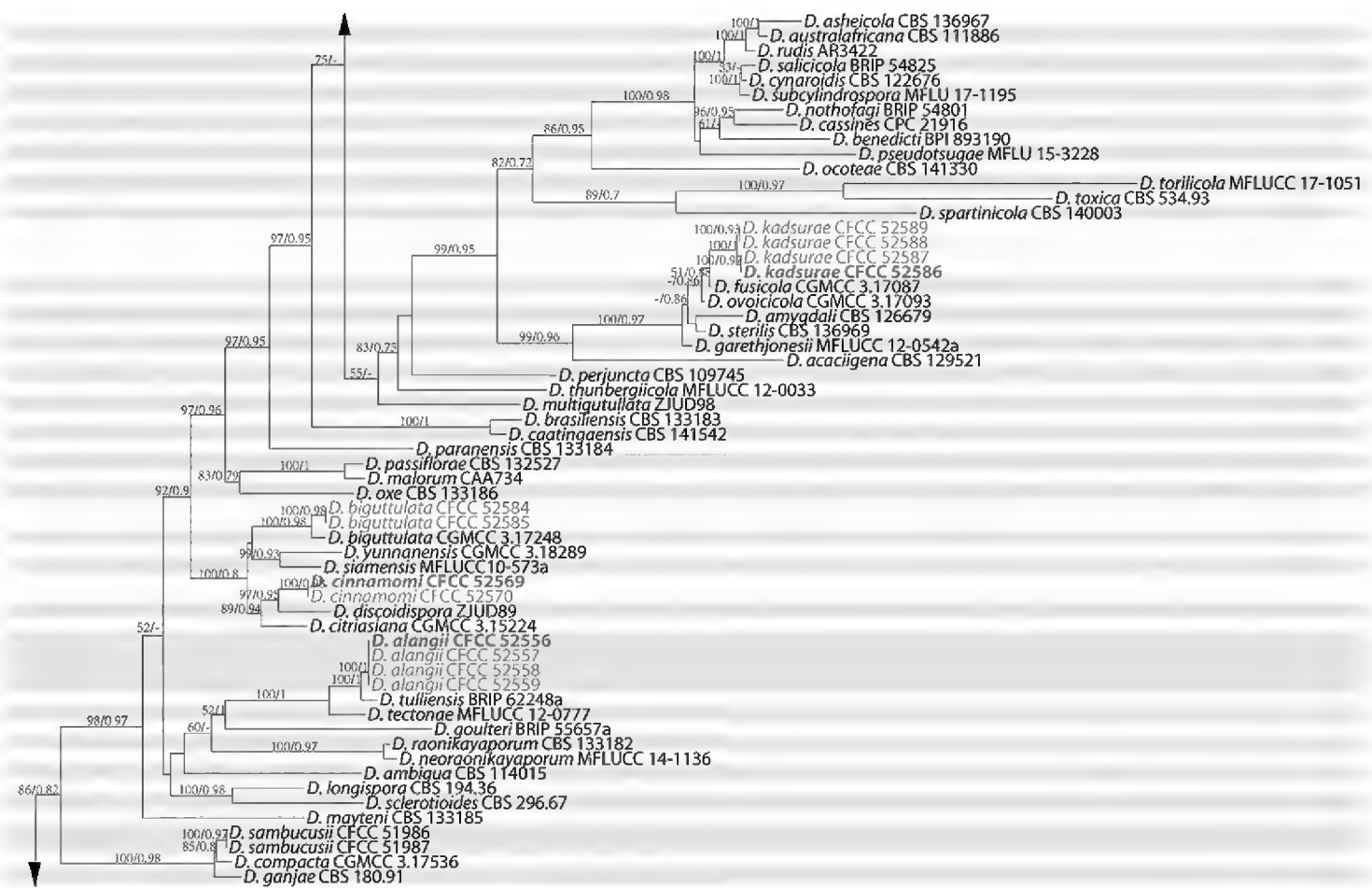


Figure I. Continued.

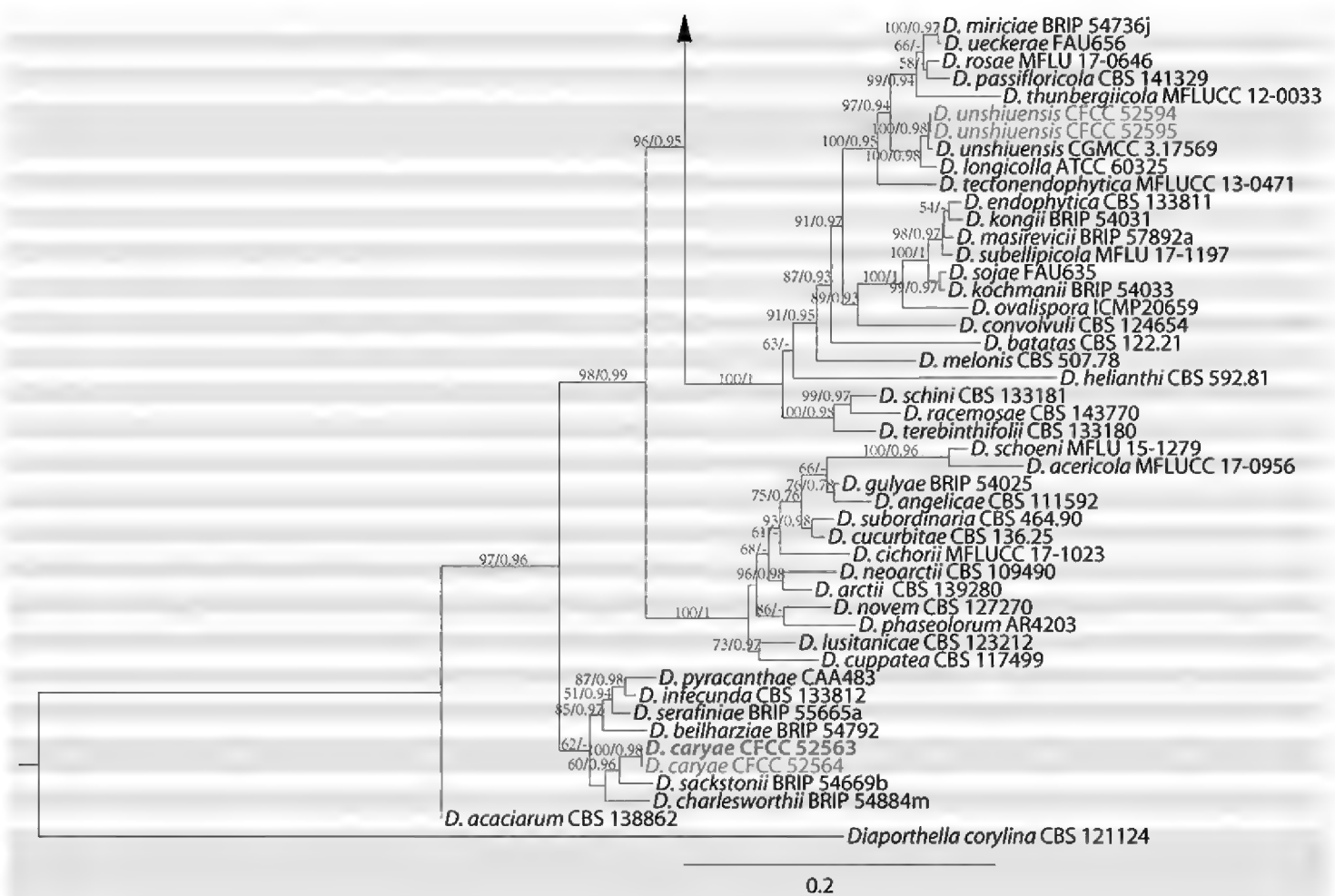


Figure I. Continued.

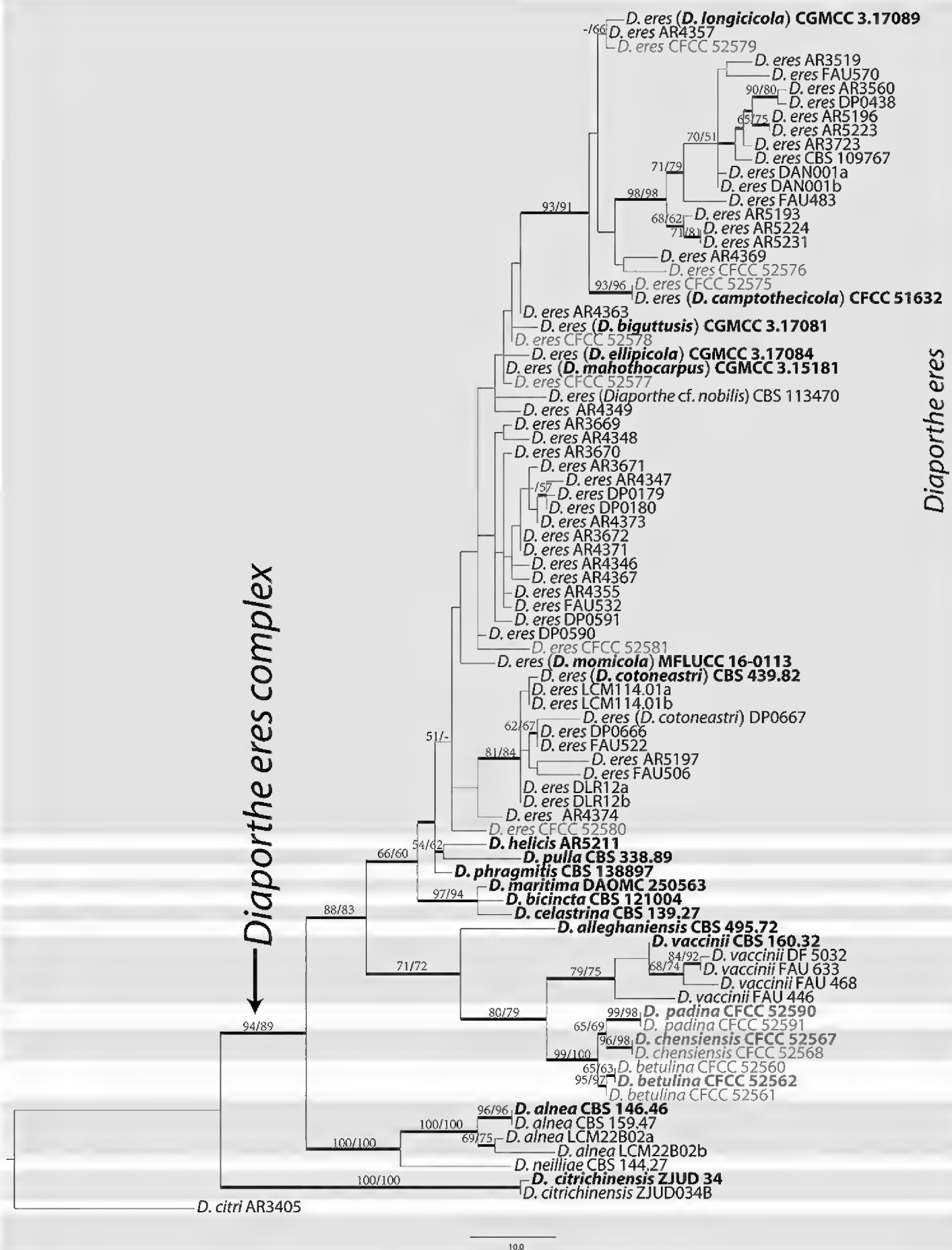


Figure 2. Phylogram of *Diaporthe eres* complex based on combined *cal*, *tef1* and *tub2*. Values above the branches indicate maximum parsimony bootstrap (left, MP BP ≥ 50%) and maximum likelihood bootstrap (right, ML BP ≥ 50%). Values below branches represent posterior probabilities (BI PP ≥ 0.70) from Bayesian inference. The tree is rooted with *Diaporthe citri*. Strains in the current study are in blue. The ex-type/ex-epitype culture is in bold.

Taxonomy

Diaporthe acerigena C.M. Tian & Q. Yang, sp. nov.

MycoBank: MB824703

Figure 3

Diagnosis. *Diaporthe acerigena* can be distinguished from the phylogenetically closely related species *D. oraccinii* in larger alpha conidia.

Holotype. CHINA. Shaanxi Province: Qinling Mountain, on symptomatic twigs of *Acer tataricum*, 27 June 2017, N. Jiang (holotype: BJFC-S1466; ex-type culture: CFCC 52554).

Etymology. Named after the host genus on which it was collected, *Acer*.

Description. On PDA: Conidiomata pycnidial, globose, solitary or aggregated, deeply embedded in the medium, erumpent, dark brown to black, 185–270 µm diam, whitish translucent to cream conidial drops exuding from the ostioles. Conidiophores 14.5–17 × 1.4–2.9 µm, cylindrical, hyaline, phialidic, branched, straight to sinuous. Alpha conidia 7–10 × 2.1–2.9 µm (av. = 8.6 × 2.5 µm, n = 30), aseptate, hyaline, ellipsoidal, rounded at one end, slightly apex at the other end, usually with two-guttulate. Beta conidia not observed.

Culture characters. Cultures incubated on PDA at 25 °C in darkness. Colony at first white, becoming dark brown in the centre with age. Aerial mycelium white, dense, fluffy, with cream conidial drops exuding from the ostioles.

Additional specimens examined. CHINA. Shaanxi Province: Qinling Mountain, on symptomatic twigs of *Acer tataricum*, 27 June 2017, N. Jiang, living culture CFCC 52555 (BJFC-S1467).

Notes. Two strains representing *D. acerigena* cluster in a well-supported clade and appear most closely related to *D. oraccinii*. *Diaporthe acerigena* can be distinguished from *D. oraccinii* based on ITS, *his3*, *tef1* and *tub2* loci (5/469 in ITS, 8/429 in *his3*, 8/326 in *tef1* and 5/358 in *tub2*). Morphologically, *D. acerigena* differs from *D. oraccinii* in the longer and larger alpha conidia (8.6 × 2.5 vs. 6.6 × 1.9 µm) (Gao et al. 2016).

Diaporthe alangii C.M. Tian & Q. Yang, sp. nov.

MycoBank: MB824704

Figure 4

Diagnosis. *Diaporthe alangii* can be distinguished from the phylogenetically closely related species *D. tectonae* and *D. tulliensis* by the size of conidiophores and alpha conidia.

Holotype. CHINA. Zhejiang Province: Tianmu Mountain, on symptomatic branches of *Alangium kurzii*, 19 Apr. 2017, Q. Yang (holotype: BJFC-S1468; ex-type culture: CFCC 52556).

Etymology. Named after the host genus on which it was collected, *Alangium*.

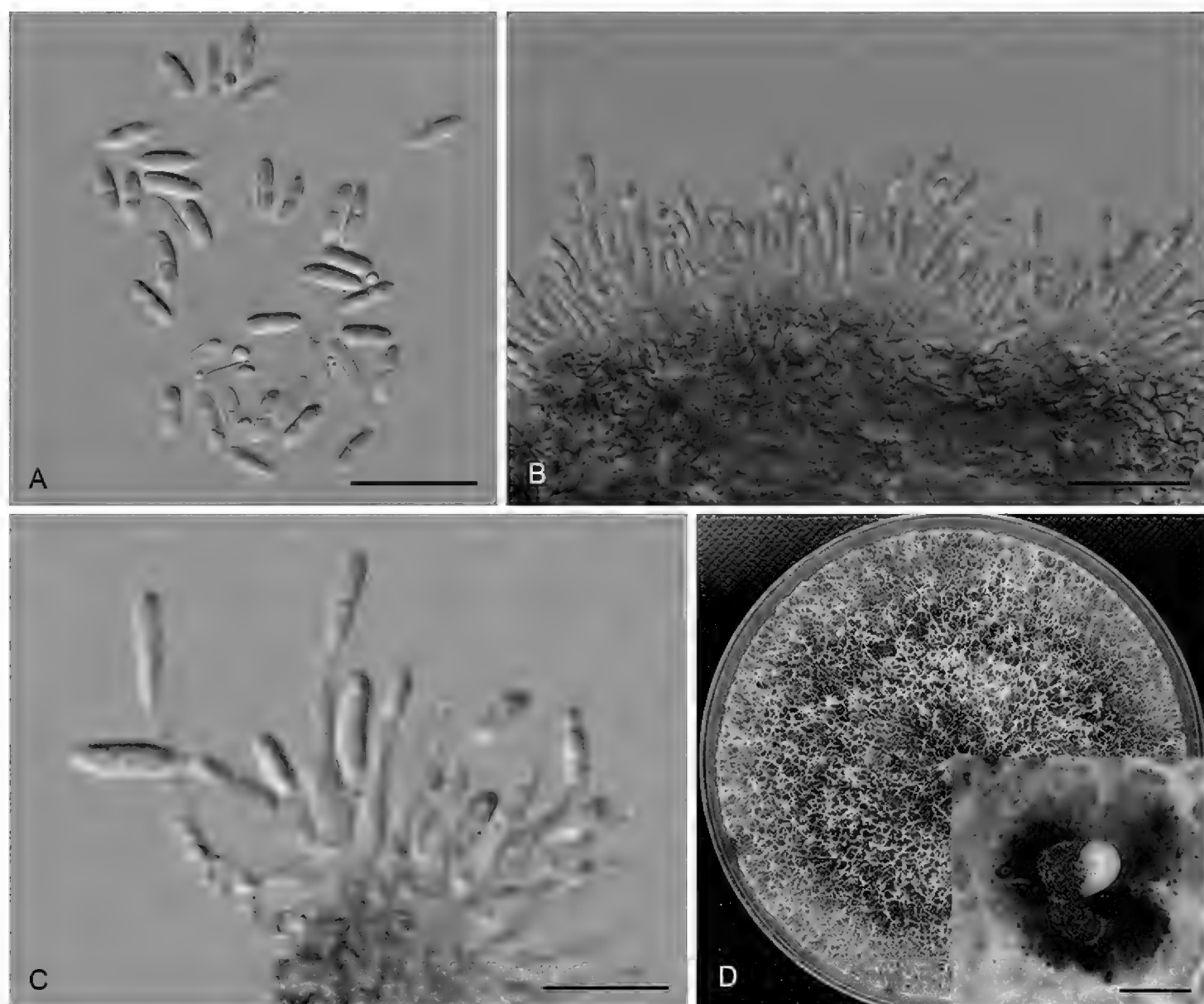


Figure 3. *Diaporthe acerigena* (CFCC 52554) **A** Alpha conidia **B–C** Conidiophores **D** Culture on PDA and conidiomata. Scale bars: 20 μm (**A–C**), 200 μm (**D**).

Description. Conidiomata pycnidial, immersed in bark, scattered, erumpent through the bark surface, discoid, with a solitary undivided locule. Ectostromatic disc black, one ostiole per disc, 135–330 μm diam. Locule circular, undivided, 290–445 μm diam. Conidiophores 6–12 \times 1.4–2 μm , cylindrical, hyaline, phialidic, unbranched, straight. Alpha conidia 6.5–8 \times 2 μm (av. = 7 \times 2 μm , $n = 30$), aseptate, hyaline, ellipsoidal, biguttulate, mostly with one end obtuse and the other acute, occasionally submedian constriction. Beta conidia not observed.

Culture characters. Cultures incubated on PDA at 25 $^{\circ}\text{C}$ in darkness. Colony initially white, producing beige pigment after 7–10 d. The colony is flat, felty with a thick texture at the centre and marginal area, with thin texture in the middle, lacking aerial mycelium, conidiomata absent.

Additional specimens examined. CHINA. Zhejiang Province: Tianmu Mountain, on symptomatic branches of *Alangium kurzii*, 19 Apr. 2017, Q. Yang, living culture CFCC 52557 (BJFC-S1469); *ibid.* living culture CFCC 52558 (BJFC-S1470); *ibid.* living culture CFCC 52559 (BJFC-S1471).

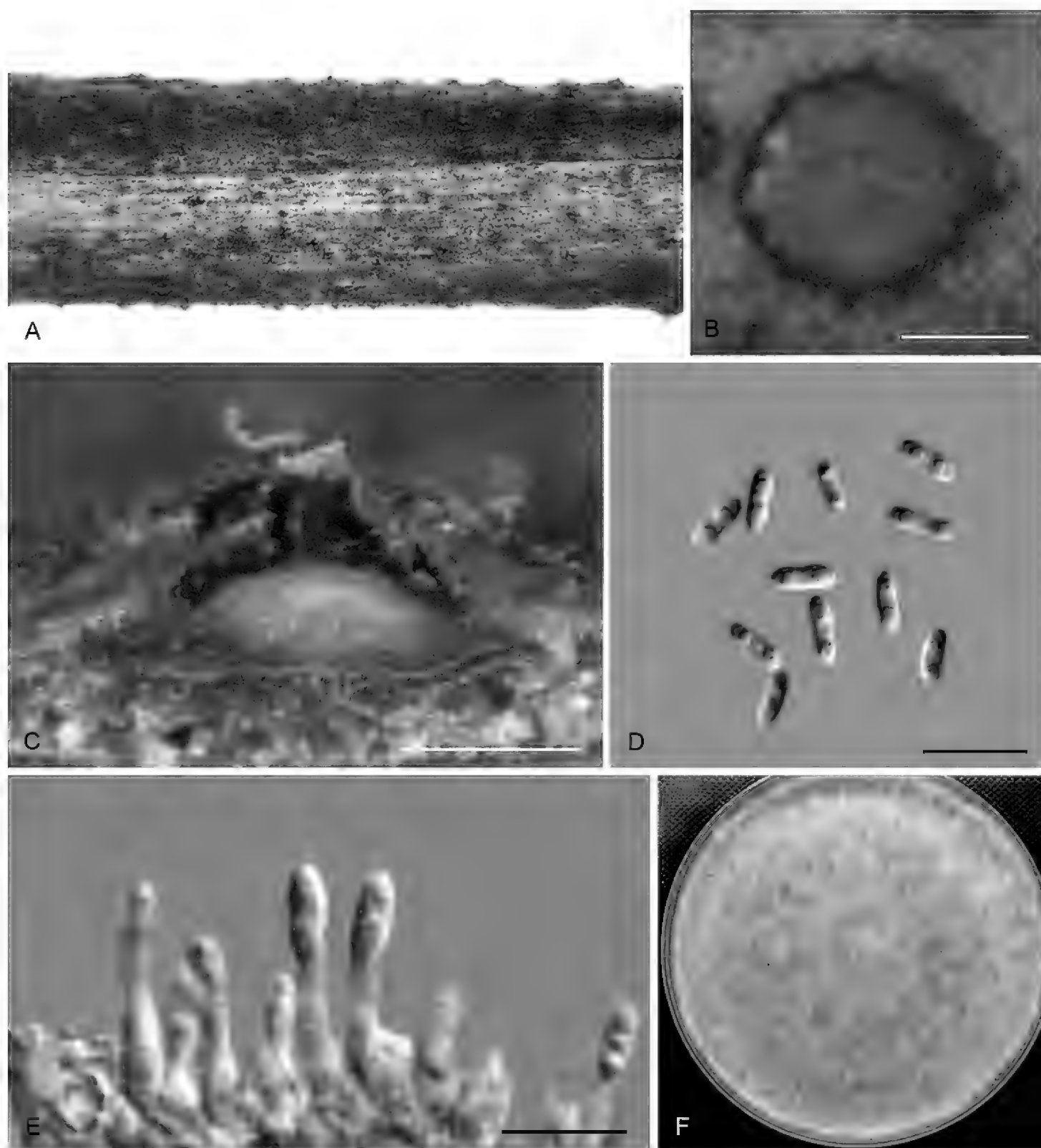


Figure 4. *Diaporthe alangii* (CFCC 52556) **A** Habit of conidiomata on branches **B** Transverse section of conidioma **C** Longitudinal section of conidioma **D** Alpha conidia **E** Conidiophores **F** Culture on PDA. Scale bars: 200 μm (**B–C**), 10 μm (**D–E**).

Notes. Four isolates clustered in a clade distinct from its closest phylogenetic neighbour, *D. tectonae* and *D. tulliensis*. *Diaporthe alangii* can be distinguished from *D. tectonae* in *cal*, *tef1* and *tub2* loci (6/458 in *cal*, 4/308 in *tef1* and 11/407 in *tub2*); from *D. tulliensis* in ITS, *tef1* and *tub2* loci (6/462 in ITS, 8/308 in *tef1* and 10/701 in *tub2*). Morphologically, *D. alangii* differs from *D. tectonae* in shorter conidiophores (6–12 vs. 11–18 μm) and longer alpha conidia (6.5–8 vs. 5.5–6 μm); from *D. tulliensis* in shorter conidiophores (6–12 vs. 15–20 μm) (Crous et al. 2015, Doilom et al. 2017).

***Diaporthe betulina* C.M. Tian & Q. Yang, sp. nov.**

MycoBank: MB824705

Figure 5

Diagnosis. *Diaporthe betulina* can be distinguished from the phylogenetically closely related species *D. betulae* in smaller locule and wider alpha conidia.

Holotype. CHINA. Heilongjiang Province: Yichun city, on symptomatic branches of *Betula platyphylla*, 27 July 2016, Q. Yang (holotype: BJFC-S1472; ex-type culture: CFCC 52562).

Etymology. Named after the host genus on which it was collected, *Betula*.

Description. Conidiomata pycnidial, conical, immersed in bark, scattered, erumpent through the bark surface, with a solitary undivided locule. Ectostromatic disc brown to black, one ostiole per disc, 290–645 µm diam. Ostiole medium black, up to the level of disc. Locule undivided, 670–905 µm diam. Conidiophores 12.5–17.5 × 1.5–2 µm, cylindrical, hyaline, phialidic, branched, straight or slightly curved. Alpha conidia hyaline, aseptate, ellipsoidal to fusiform, 0–2-guttulate, sometimes acute at both ends, 8–10 × 2.5–3 µm (av. = 9 × 2.6 µm, n = 30). Beta conidia hyaline, aseptate, filiform, straight or hamate, eguttulate, base subtruncate, tapering towards one apex, 26–32.5 × 1 µm (av. = 30 × 1 µm, n = 30).

Culture characters. Cultures incubated on PDA at 25 °C in darkness. Colony flat with white felty aerial mycelium, turning white to dark brown aerial mycelium, conidiomata irregularly distributed on the agar surface.

Additional specimens examined. CHINA. Heilongjiang Province: Yichun city, on symptomatic branches of *Betula albo-sinensis*, 27 July 2016, Q. Yang, living culture CFCC 52560 (BJFC-S1473); on symptomatic branches of *Betula costata*, 27 July 2016, Q. Yang, living culture CFCC 52561 (BJFC-S1474).

Notes. *Diaporthe betulina* was isolated from *Betula* spp. cankers in Heilongjiang Province. Three strains representing *D. betulina* cluster in a well-supported clade and appear most closely related to *D. betulae*, which was also isolated from *Betula platyphylla* in Sichuan Province (Du et al. 2016). *Diaporthe betulina* can be distinguished based on ITS, *his3*, *tef1* and *tub2* loci from *D. betulae* (11/461 in ITS, 9/453 in *his3*, 12/336 in *tef1* and 7/695 in *tub2*). Morphologically, *D. betulina* differs from *D. betulae* in smaller locule (470–945 vs. 600–1250 µm) and wider alpha conidia (3–4 vs. 2.5–3 µm) (Du et al. 2016).

***Diaporthe biguttulata* F. Huang, K.D. Hyde & H.Y. Li, 2015**

Figure 6

Description. Conidiomata pycnidial, immersed in bark, scattered, erumpent through the bark surface, discoid, with a single locule. Ectostromatic disc dark brown, one ostiole per disc, 160–320 µm diam. Locule undivided, 235–350 µm diam. Conidiophores 8.5–11 × 1.5 µm, cylindrical, hyaline, branched, straight or slightly curved, tapering

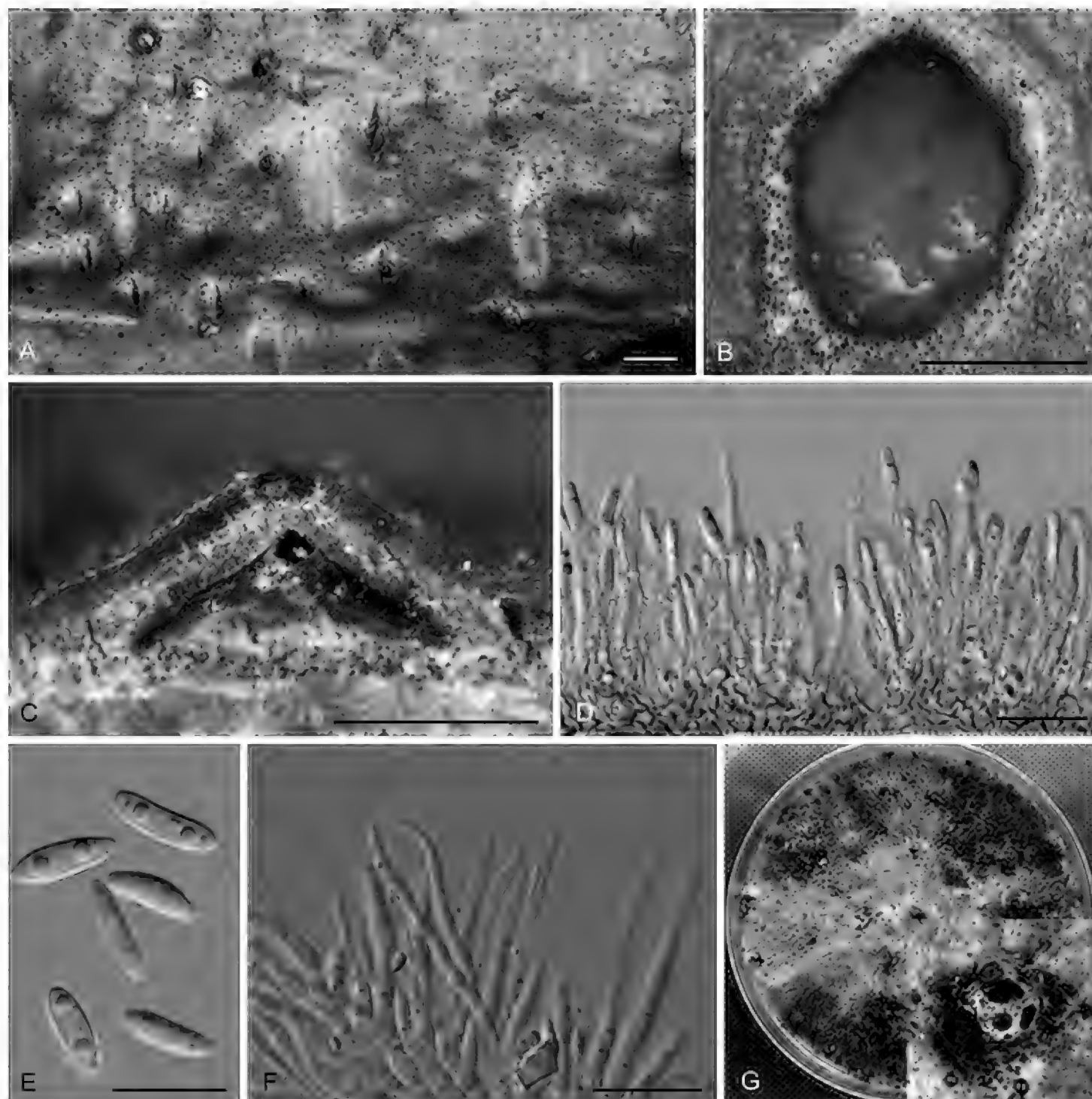


Figure 5. *Diaporthe betulina* (CFCC 52562) **A** Habit of conidiomata on branches **B** Transverse section of conidioma **C** Longitudinal section of conidioma **D** Conidiophores **E** Alpha conidia **F** Beta conidia **G** Culture on PDA and conidiomata. Scale bars: 500 μm (**A–C**), 10 μm (**D–F**).

towards the apex. Alpha conidia hyaline, aseptate, ellipsoidal to oval, 2-guttulate, usually rounded at both ends, occasionally with one end acute, $7\text{--}8.5 \times 1.5\text{--}2 \mu\text{m}$ (av. = $6.5 \times 2.6 \mu\text{m}$, $n = 30$). Beta conidia not observed.

Culture characters. Cultures incubated on PDA at 25 °C in darkness. Colony originally flat with white aerial mycelium, becoming pale grey, with dense aerial mycelium in the centre and sparse aerial mycelium at the marginal area, conidiomata absent.

Specimens examined. CHINA. Zhejiang Province: Tianmu Mountain, on symptomatic branches of *Juglans regia*, 20 Apr. 2017, Q. Yang, living culture CFCC 52584 and CFCC 52585 (BJFC-S1504).

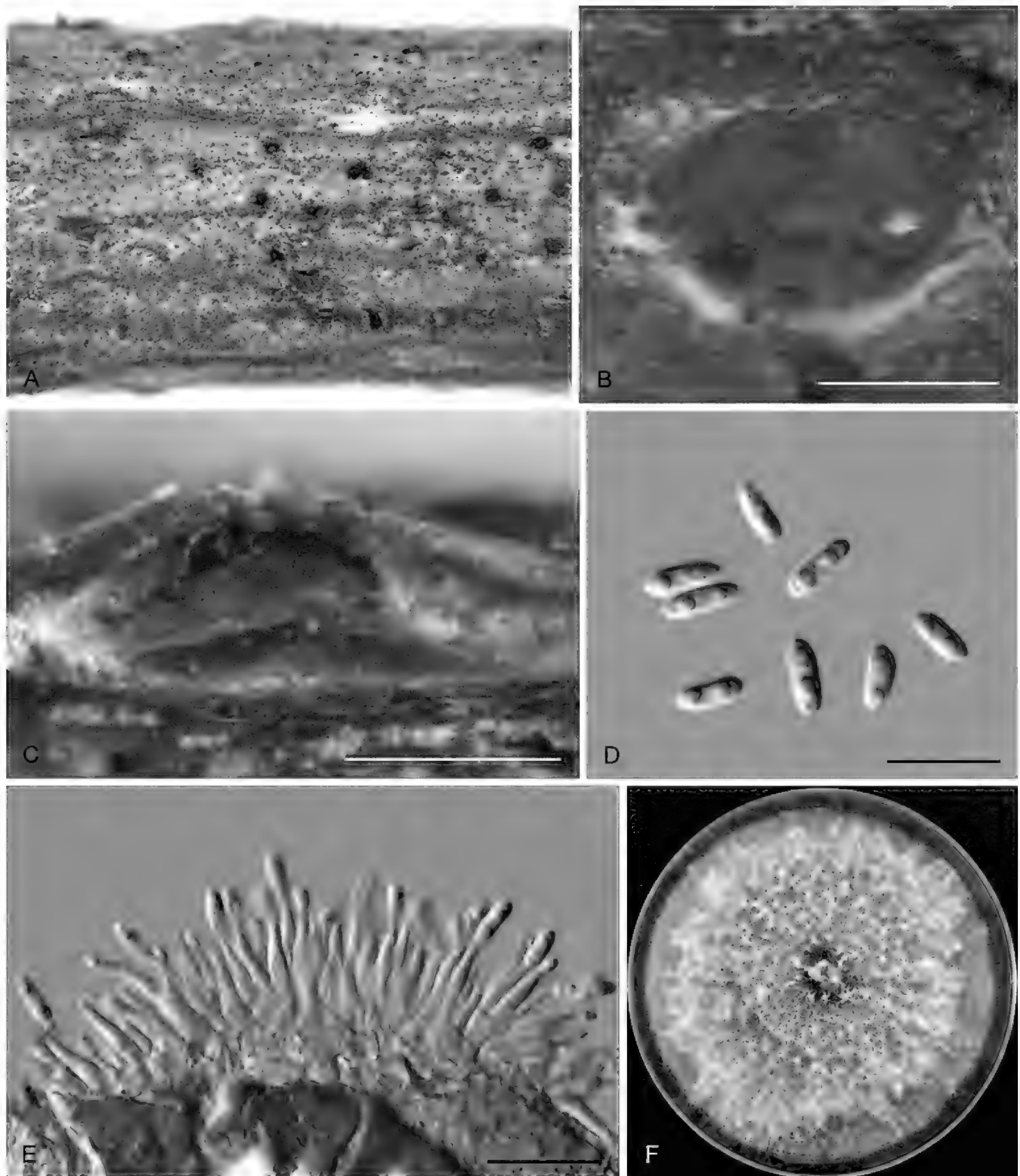


Figure 6. *Diaporthe biguttulata* (CFCC 52584) **A** Habit of conidiomata on branches **B** Transverse section of conidioma **C** Longitudinal section of conidioma **D** Alpha conidia **E** Conidiophores **F** Culture on PDA. Scale bars: 200 μm (**B–C**), 10 μm (**D–E**).

Notes. *Diaporthe biguttulata* was originally described from a healthy branch of *Citrus limon* in Yunnan Province, China (Huang et al. 2015). In the present study, two isolates (CFCC 52584 and CFCC 52585) from symptomatic branches of *Juglans regia* were congruent with *D. biguttulata* based on morphology and DNA sequences data (Fig. 1). We therefore describe *D. biguttulata* as a known species for this clade.

***Diaporthe caryae* C.M. Tian & Q. Yang, sp. nov.**

MycoBank: MB824706

Figure 7

Diagnosis. *Diaporthe caryae* differs from its closest phylogenetic neighbour, *D. charlesworthii* and *D. sackstonii*, in ITS, *tef1* and *tub2* loci based on the alignments deposited in TreeBASE.

Holotype. CHINA. Jiangsu Province: Nanjing city, on symptomatic twigs of *Carya illinoensis*, 10 Nov. 2015, Q. Yang (holotype: BJFC-S1476; ex-type culture: CFCC 52563).

Etymology. Named after the host genus on which it was collected, *Carya*.

Description. Conidiomata pycnidial, immersed in bark, scattered, slightly erumpent through the bark surface, nearly flat, discoid, with a solitary undivided locule. Ectostromatic disc brown to black, one ostiole per disc. Locule undivided, 310–325 µm diam. Conidiophores 7–11 × 1.4–2.2 µm, cylindrical, phialidic, unbranched, sometimes inflated. Alpha conidia hyaline, aseptate, ellipsoidal or fusiform, eguttulate, obtuse at both ends, 7–8.5 × 2.1–2.5 µm (av. = 8 × 2.3 µm, n = 30). Beta conidia hyaline, aseptate, filiform, straight or hamate, eguttulate, base subtruncate, tapering towards one apex, 15.5–34 × 1.1–1.4 µm (av. = 27.5 × 1.2 µm, n = 30).

Culture characters. Cultures incubated on PDA at 25 °C in darkness. Colony at first flat with white felty mycelium, becoming black in the centre and black at the marginal area with age, conidiomata not observed.

Additional specimens examined. CHINA. Jiangsu Province: Nanjing city, on symptomatic twigs of *Carya illinoensis*, 10 Nov. 2015, Q. Yang, living culture CFCC 52564 (BJFC-S1477).

Notes. Two strains representing *D. caryae* cluster in a well-supported clade and appear closely related to *D. charlesworthii* and *D. sackstonii*. *Diaporthe caryae* can be distinguished based on ITS, *tef1* and *tub2* loci from *D. charlesworthii* (50/468 in ITS, 107/338 in *tef1* and 90/707 in *tub2*); from *D. sackstonii* (4/440 in ITS, 13/340 in *tef1* and 23/701 in *tub2*). Morphologically, *D. caryae* can be distinguished from *D. charlesworthii* by its shorter conidiophores (7–11 vs. 15–35 µm); from *D. sackstonii* by its longer alpha conidia (7–8.5 vs. 6–7 µm) (Thompson et al. 2015).

***Diaporthe cercidis* C.M. Tian & Q. Yang, sp. nov.**

MycoBank: MB824707

Figure 8

Diagnosis. *Diaporthe cercidis* can be distinguished from the phylogenetically closely related species *D. pescicola* in larger alpha conidia.

Holotype. CHINA. Jiangsu Province: Nanjing city, on twigs and branches of *Cercis chinensis*, 11 Nov. 2015, Q. Yang (holotype: BJFC-S1478; ex-type culture: CFCC 52565).

Etymology. Named after the host genus on which it was collected, *Cercis*.

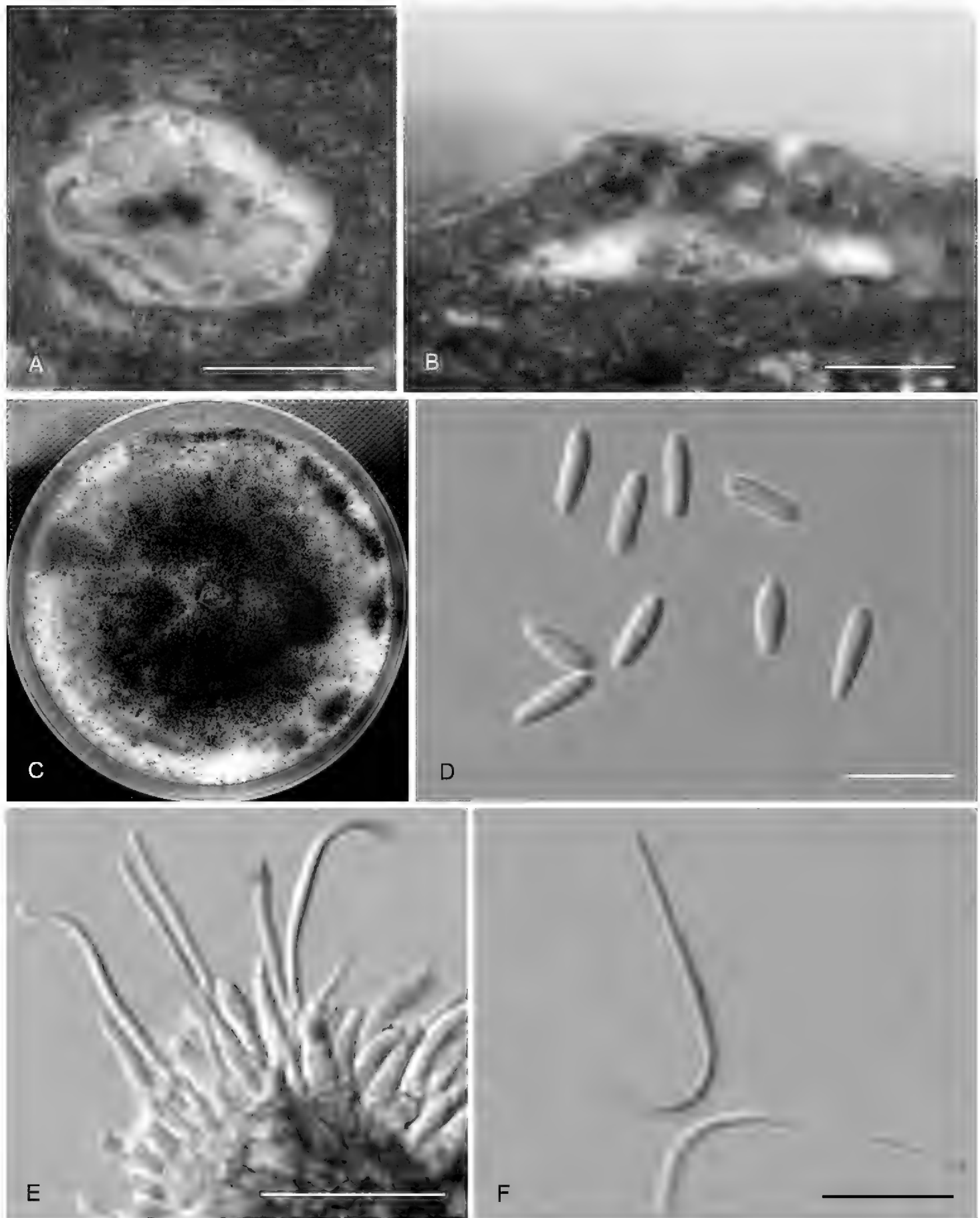


Figure 7. *Diaporthe caryae* (CFCC 52563) **A** Transverse section of conidioma **B** Longitudinal section of conidioma **C** Culture on PDA **D** Alpha conidia **E** Conidiophores **F** Beta conidia. Scale bars: 200 µm (**A**), 100 µm (**B**), 10 µm (**D, F**), 20 µm (**E**).

Description. Conidiomata pycnidial, immersed in bark, scattered, slightly erumpent through the bark surface, nearly flat, discoid, with a solitary undivided locule. Ectostromatic disc grey to brown, one ostiole per disc. Locule circular, undivided, 135–200 µm diam. Conidiophores 7–17 × 1.4–2.1 µm, phialidic, unbranched, straight or slightly curved, tapering towards the apex. Alpha conidia hyaline, aseptate, fusiform to oval, bi-

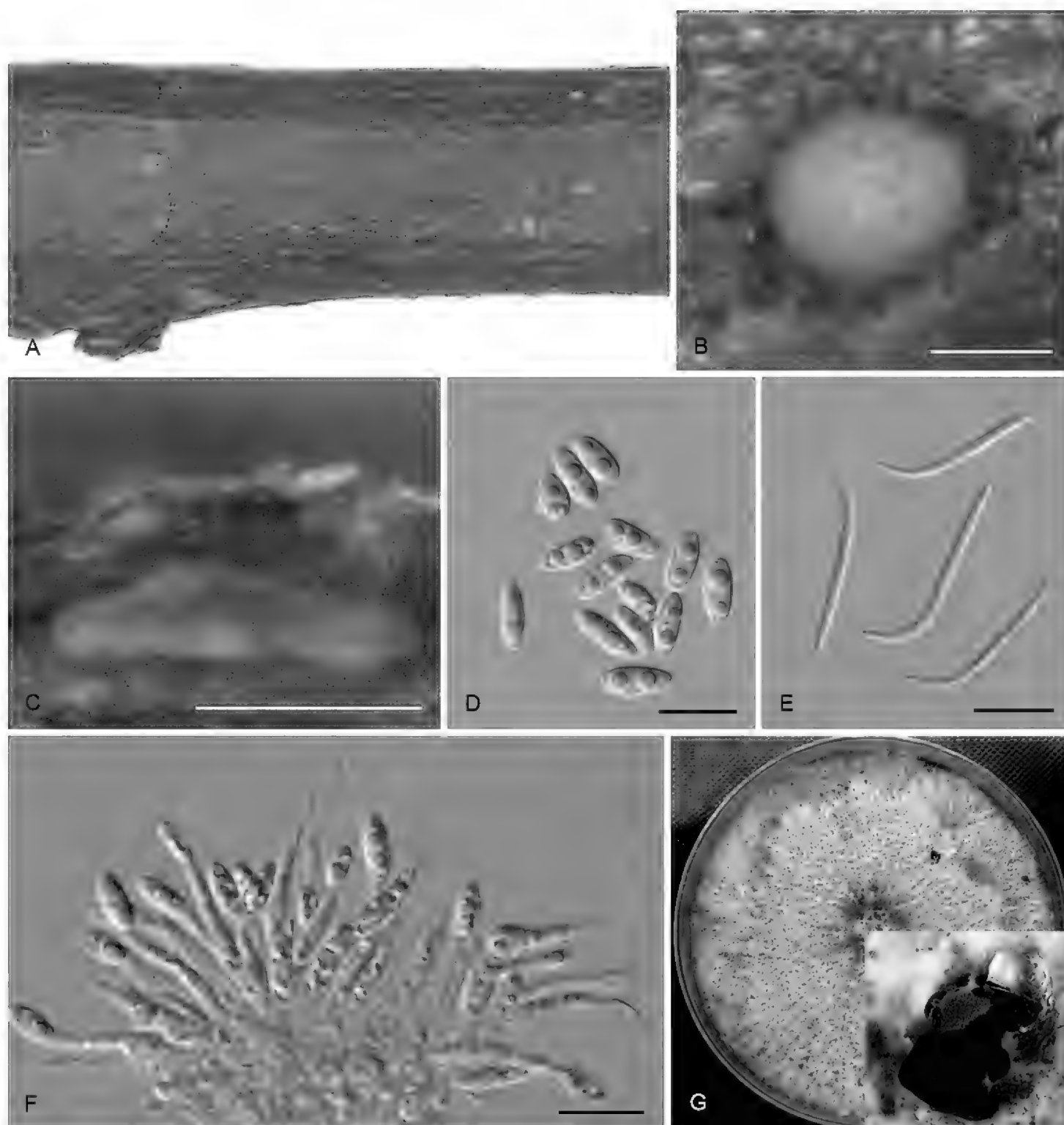


Figure 8. *Diaporthe cercidis* (CFCC 52565) **A** Habit of conidiomata on branches **B** Transverse section of conidioma **C** Longitudinal section of conidioma **D** Alpha conidia **E** Beta conidia **F** Conidiophores **G** Culture on PDA and conidiomata. Scale bars: 100 μm (**B–C**), 10 μm (**D–F**).

guttulate, $6.5\text{--}10 \times 3\text{--}3.5 \mu\text{m}$ (av. = $8.6 \times 3.3 \mu\text{m}$, $n = 30$). Beta conidia hyaline, aseptate, filiform, straight or hamate, eguttulate, $20\text{--}28.5 \times 1\text{--}1.3 \mu\text{m}$ (av. = $25.5 \times 1.2 \mu\text{m}$, $n = 30$).

Culture characters. Cultures incubated on PDA at 25 °C in darkness showed colony at first white, becoming pale brown with yellowish dots with age, flat, with dense and felted mycelium, with visible solitary or aggregated conidiomata at maturity.

Additional specimens examined. CHINA. Jiangsu Province: Yangzhou city, on twigs and branches of *Ginkgo biloba*, 11 Nov. 2015, N. Jiang, living culture CFCC 52566 (BJFC-S1479).

Notes. *Diaporthe cercidis* is distinguished from *D. pescicola* in the ITS, *cal* and *tef1* loci (13/458 in ITS, 47/442 in *cal* and 6/328 in *tef1*). Morphologically, *D. cercidis* dif-

fers from *D. pescicola* in shorter conidiophores (7–17 vs. 21–35 µm) and larger alpha conidia (6.5–10 × 3–3.5 vs. 6–8.5 × 2–3 µm) (Dissanayake et al. 2017a).

***Diaporthe chensiensis* C.M. Tian & Q. Yang, sp. nov.**

MycoBank: MB824708

Figure 9

Diagnosis. *Diaporthe chensiensis* differs from its closest phylogenetic neighbour, *D. vaccinii*, in ITS, *cal*, *his3* and *tef1* loci based on the alignments deposited in TreeBASE.

Holotype. CHINA. Shaanxi Province: Ningshan County, Huoditang forest farm, on symptomatic twigs of *Abies chensiensis*, 5 July 2017, Q. Yang (holotype: BJFC-S1480; ex-type culture: CFCC 52567).

Etymology. Named after the host species on which it was collected, *chensiensis*.

Description. Conidiomata pycnidial, immersed in bark, scattered, slightly erumpent through the bark surface, discoid, with a single locule. Ectostromatic disc white to brown, one ostiole per disc, 200–325 µm diam. Locule undivided, 385–540 µm diam. Conidiophores 8.5–13 × 2–3 µm, cylindrical, hyaline, phialidic, unbranched, straight or slightly curved, tapering towards the apex. Alpha conidia hyaline, aseptate, smooth, ellipsoidal, biguttulate, rounded at both ends, 6.5–11 × 2–2.2 µm (av. = 8.5 × 2.1 µm, n = 30). Beta conidia present on the host, hyaline, eguttulate, smooth, filiform, hamate, 21–28.5 × 0.8–1.1 µm (av. = 25 × 1 µm, n = 30).

Culture characters. Cultures incubated on PDA at 25 °C in darkness. Colony originally flat with white felted aerial mycelium, becoming light brown mycelium due to pigment formation, conidiomata irregularly distributed over agar surface, with yellowish conidial drops exuding from the ostioles.

Additional specimens examined. CHINA. Shaanxi Province: Ningshan County, Huoditang forest farm, on symptomatic twigs of *Abies chensiensis*, 5 July 2017, Q. Yang, living culture CFCC 52568 (BJFC-S1481).

Notes. *Diaporthe chensiensis* occurs in an independent clade (Fig. 1) and is phylogenetically distinct from *D. vaccinii*. *Diaporthe chensiensis* can be distinguished from *D. vaccinii* by 57 nucleotides in concatenated alignment, in which 14 were distinct in the ITS region, 13 in the *cal* region, 10 in the *his3* region, 15 in the *tef1* region and 15 in the *tub2* region. Although this species belongs to the *D. eres* complex, it is, however, distinct from the known species within the complex (Fig. 2).

***Diaporthe cinnamomi* C.M. Tian & Q. Yang, sp. nov.**

MycoBank: MB824709

Figure 10

Diagnosis. *Diaporthe cinnamomi* differs from its closest phylogenetic species *D. discoidispora* in ITS, *his3* and *tef1* loci based on the alignments deposited in TreeBASE.

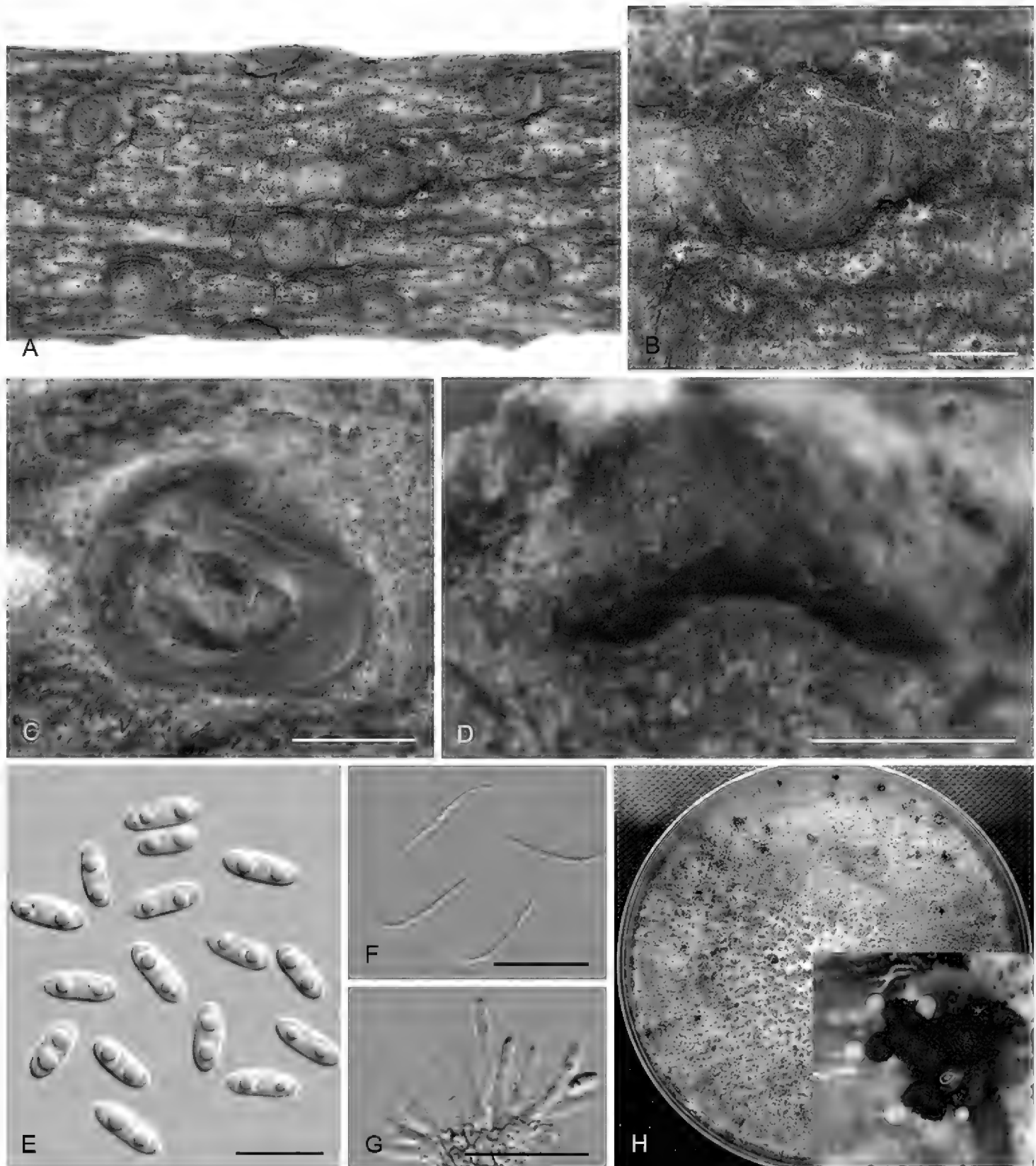


Figure 9. *Diaporthe chensiensis* (CFCC 52567) **A–B** Habit of conidiomata on branches **C** Transverse section of conidioma **D** Longitudinal section of conidioma **E** Alpha conidia **F** Beta conidia **G** Conidiophores **H** Culture on PDA and conidiomata. Scale bars: 500 μm (**B**), 200 μm (**C–D**), 10 μm (**E**), 20 μm (**F**).

Holotype. CHINA. Zhejiang Province: Linan city, on symptomatic twigs of *Cinnamomum* sp., 22 Apr. 2017, Q. Yang (holotype: BJFC-S1482; ex-type culture: CFCC 52569).

Etymology. Named after the host genus on which it was collected, *Cinnamomum*.

Description. On PDA: Conidiomata pycnidial, globose, solitary or aggregated, deeply embedded in the substrate, erumpent, dark brown to black, 170–235 μm diam., whitish translucent to cream conidial drops exuding from the ostioles. Conidiophores

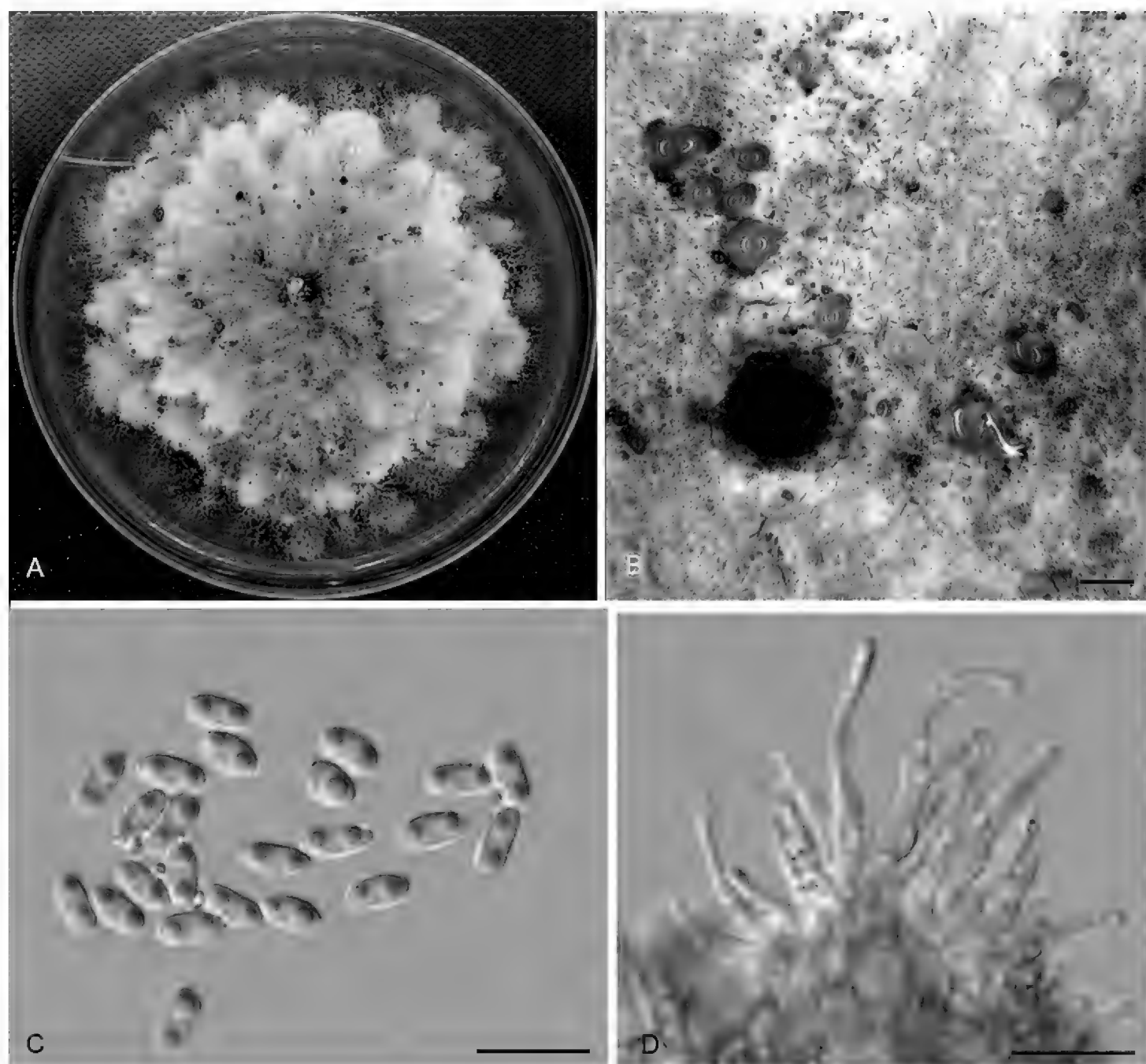


Figure 10. *Diaporthe cinnamomi* (CFCC 52569) **A** Culture on PDA **B** Conidiomata **C** Alpha conidia **D** Conidiophores. Scale bars: 200 μm (**B**), 10 μm (**C–D**).

11–25 \times 1.5–2 μm , cylindrical, hyaline, branched, straight or curved, tapering towards the apex. Alpha conidia hyaline, aseptate, ellipsoidal to oval, biguttulate, rounded at both ends, 5–7 \times 2.5–3 μm (av. = 6 \times 2.9 μm , $n = 30$). Beta conidia not observed.

Culture characters. Cultures incubated on PDA at 25 $^{\circ}\text{C}$ in darkness showed colony originally flat with white felty mycelium, developing petaloid mycelium after 7–10 d and turning yellowish at the centre and brownish at the marginal area after 15 d. Conidiomata erumpent at maturity.

Additional material examined. CHINA. Zhejiang Province: Linan city, on symptomatic twigs of *Cinnamomum* sp., 22 Apr. 2017, Q. Yang, living culture CFCC 52570 (BJFC-S1483).

Notes. *Diaporthe cinnamomi* comprises strains CFCC 52569 and CFCC 52570 closely related to *D. discoidispora* in the combined phylogenetic tree (Fig. 1). *Diaporthe cinnamomi* can be distinguished based on ITS, *his3* and *tef1* loci from *D. discoidispora* (4/460 in ITS, 17/448 in *his3* and 38/339 in *tef1*).

***Diaporthe conica* C.M. Tian & Q. Yang, sp. nov.**

MycoBank: MB824710

Figure 11

Diagnosis. *Diaporthe conica* is phylogenetically and morphologically distinct from *D. rostrata*, in smaller locule and alpha conidia.

Holotype. CHINA. Zhejiang Province: Tianmu Mountain, on symptomatic branches of *Alangium chinense*, 20 Apr. 2017, Q. Yang (holotype: BJFC-S1484; ex-type culture: CFCC 52571).

Etymology. Named after the conical conidiomata.

Description. Conidiomata pycnidial, 420–580 µm diam., solitary and with single necks erumpent through the host bark. Tissue around the neck is conical. Locule oval, undivided, 385–435 µm diam. Conidiophores reduced to conidiogenous cells. Conidiogenous cells unbranched, straight or sinuous, apical or base sometimes swelling, 19–23.5 × 2.8 µm. Alpha conidia hyaline, aseptate, ellipsoidal, biguttulate, 5.5–7 × 2.3–3 µm (av. = 6.5 × 2.6 µm, n = 30). Beta conidia not observed.

Culture characters. Cultures incubated on PDA at 25 °C in darkness. Colony white to yellowish, with dense and felted mycelium, lacking aerial mycelium, with maize-coloured conidial drops exuding from the ostioles.

Additional material examined. CHINA. Zhejiang Province: Tianmu Mountain, on symptomatic branches of *Alangium chinense*, 20 Apr. 2017, Q. Yang, living culture CFCC 52572 (BJFC-S1485); ibid. living culture CFCC 52573 (BJFC-S1486); ibid. living culture CFCC 52574 (BJFC-S1487).

Notes. Four isolates clustered in a clade distinct from further *Diaporthe* species based on DNA sequence data. Morphologically, this species is characterised by conical conidiomata, which is similar with *D. rostrata* from *Juglans mandshurica*. However, *D. conica* differs from *D. rostrata* by having smaller locule and alpha conidia (310–385 vs. 620–1100 µm in locule; 5.5–7 × 2.3–3 vs. 8.5–11.5 × 4–5 µm in alpha conidia) (Fan et al. 2015).

***Diaporthe eres* Nitschke, 1870**

Figure 12

= *Diaporthe biguttusis* Y.H. Gao & L. Cai, 2015.

= *Diaporthe camptothecicola* C.M. Tian & Qin Yang, 2017.

= *Diaporthe ellipicola* Y.H. Gao & L. Cai, 2015.

= *Diaporthe longicicola* Y.H. Gao & L. Cai, 2015

= *Diaporthe mahothocarpus* (Y.H. Gao, W. Sun & L. Cai) Y.H. Gao & L. Cai, 2015.

= *Diaporthe momicola* Dissan., J.Y. Yan, Xing H. Li & K.D. Hyde, 2017.

Description. Conidiomata pycnidial, immersed in bark, erumpent through the bark surface, serried, with a single locule. Ectostromatic disc obviously, brown to black,

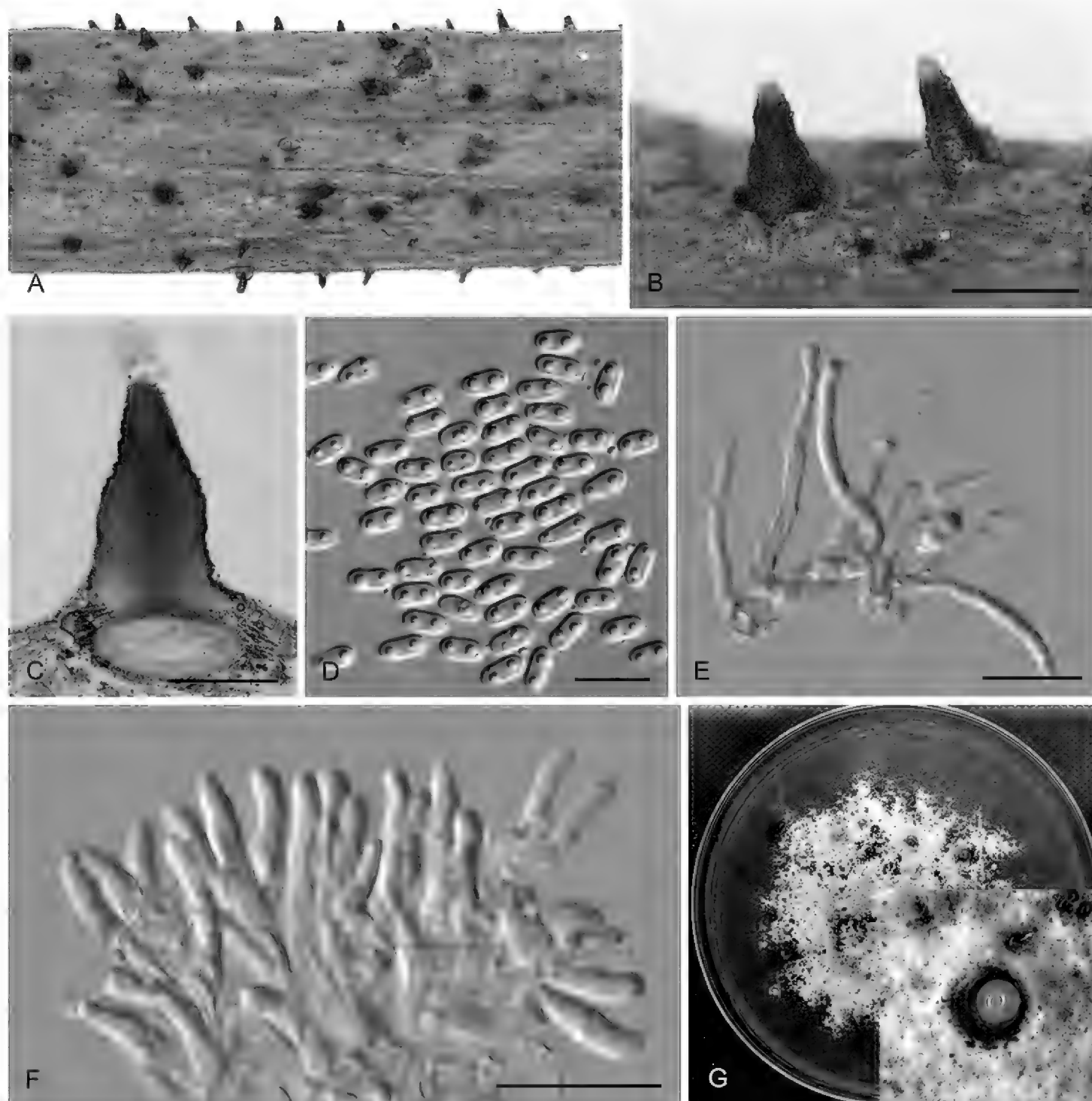


Figure 11. *Diaporthe conica* (CFCC 52571) **A–B** Habit of conidiomata on branches **C** Longitudinal section of conidioma **D** Alpha conidia **E–F** Conidiophores **G** Culture on PDA and conidiomata. Scale bars: 300 μm (**B–C**), 10 μm (**D–F**).

with one ostiole per disc, 245–572 μm diam. Ostiole medium black, up to the level of disc. Locule circular, undivided, 335–450 μm diam. Conidiophores 10.5–19 \times 1–1.5 μm , cylindrical, hyaline, unbranched, straight or slightly sinuous. Conidiogenous cells phialidic, cylindrical, terminal. Alpha conidia hyaline, aseptate, ellipsoidal to lanceolate, one guttulate at each end, 6–7.5 \times 1.5–2.5 μm (av. = 6.5 \times 2 μm , $n = 30$). Beta conidia not observed.

Culture characters. Cultures on PDA incubated at 25 $^{\circ}\text{C}$ in darkness. Colony with white felty aerial mycelium, becoming white felted aerial mycelium in the centre and grey-brown mycelium at the marginal area, conidiomata irregularly distributed over agar surface.

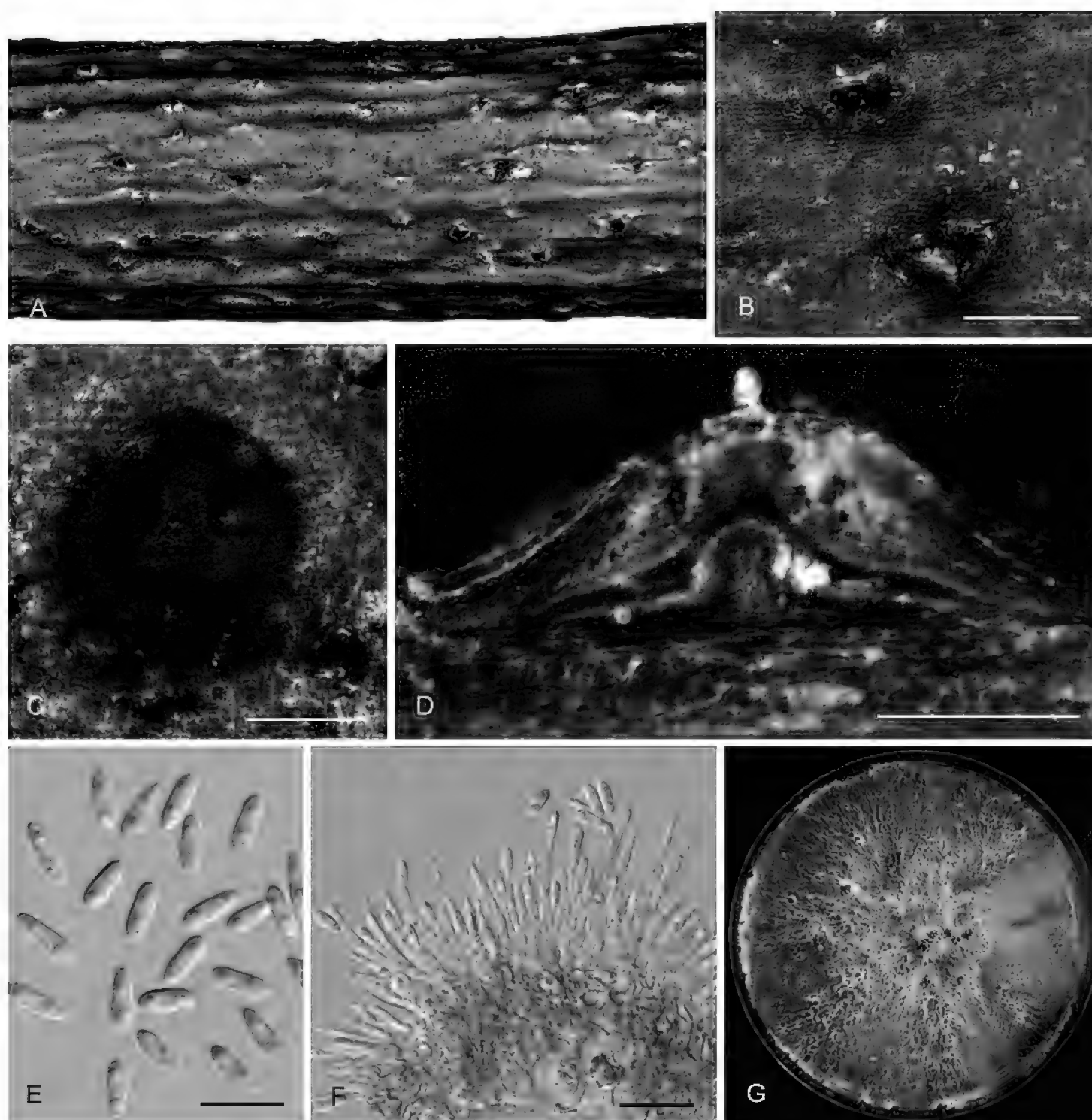


Figure 12. *Diaporthe eres* (CFCC 52575) **A–B** Habit of conidiomata on branches **C** Transverse section of conidioma **D** Longitudinal section of conidioma **E** Alpha conidia **F** Conidiophores **G** Culture on PDA and conidiomata. Scale bars: 500 μm (**B**), 200 μm (**C–D**), 10 μm (**E–F**).

Specimens examined. CHINA. Beijing: Pinggu district, on symptomatic branches of *Castanea mollissima*, 1 Nov. 2016, N. Jiang, living culture CFCC 52576 (BJFC-S1489); *ibid.* living culture CFCC 52577 (BJFC-S1490). Heilongjiang Province: Liangshui Nature Reserve, on symptomatic twigs of *Acanthopanax senticosus*, 29 July 2016, Q. Yang, living culture CFCC 52580 (BJFC-S1493). Heilongjiang Province: Harbin city, Botanical garden, on symptomatic twigs of *Sorbus* sp., 2 Aug. 2016, Q. Yang, living culture CFCC 52575 (BJFC-S1488). Shaanxi Province: Zhashui County, on symptomatic branches of *Juglans regia*, 29 July 2016, Q. Yang, living culture CFCC 52579 (BJFC-S1492). Zhejiang Province: Yangzhou city, on symptomatic twigs of

Melia azedarace, 8 July 2017, N. Jiang, living culture CFCC 52578 (BJFC-S1491). Zhejiang Province: Tianmu Mountain, on symptomatic twigs of *Rhododendron simsii*, 20 Apr. 2017, Q. Yang, living culture CFCC 52581 (BJFC-S1494).

Notes. *Diaporthe eres*, the type species of the genus, was described by Nitschke (1870) on *Ulmus* sp. collected in Germany, which has a widespread distribution and a broad host range as a pathogen, endophyte or saprobe causing leaf spots, stem cankers and diseases of woody plants (Udayanga et al. 2014b). Fan et al. (2018) indicated that *D. biguttusis*, *D. ellipicola*, *D. longicicola* and *D. mahothocarpus* should be treated as synonyms of *D. eres* using *cal*, *tef1* and *tub2* gene regions. In this study, we extended the work presented in Fan et al. (2018) and found seven additional strains belonging to *D. eres*. Additionally, the phylogenetic tree demonstrated that *D. camptothecicola* and *D. momicola* should also be treated as synonyms of *D. eres* (Fig. 2). *Diaporthe camptothecicola* from *Camptotheca acuminata* and *D. momicola* from *Prunus persica* are described and illustrated based on the combined ITS, *cal*, *his3*, *tef1* and *tub2* regions (Dissanayake et al. 2017a, Yang et al. 2017c). Both of the two species are embedded in the *D. eres* complex. However, ITS analysis resulted in an unresolved phylogenetic tree without definitive bootstrap at the internodes, highly discordant to the trees resulting from the other four genes (Udayanga et al. 2014b). Therefore, the ITS region was not used in the combined analysis in the current study. To further investigate this complex, a second set of four (*cal*, *his3*, *tef1* and *tub2*), three (*cal*, *tef1* and *tub2*), two (*tef1* and *tub2*) and one (*tef1*) data matrices were performed following Santos et al. (2017) and Fan et al. (2018). The results showed that the three genes analyses (*cal*, *tef1* and *tub2*) appeared to be a better species recognition (Fig. 2). When it comes to this species complex, sequences supported by Udayanga et al. (2014b) are necessary to perform a more robust phylogenetic tree, clarifying the real species boundaries in this group in the future work.

***Diaporthe fraxinicola* C.M. Tian & Q. Yang, sp. nov.**

MycoBank: MB824711

Figure 13

Diagnosis. *Diaporthe fraxinicola* can be distinguished from the closely related species *D. oraccinii* and *D. acerigena* (described above) based on ITS, *tef1* and *tub2* loci. *Diaporthe fraxinicola* differs from *D. oraccinii* in larger alpha conidia and from *D. acerigena* in wider alpha conidia.

Holotype. CHINA. Shaanxi Province: Zhashui city, Niubeiliang Reserve, on symptomatic twigs of *Fraxinus chinensis*, 7 July 2017, Q. Yang (holotype: BJFC-S1495; ex-type culture: CFCC 52582).

Etymology. Named after the host genus on which it was collected, *Fraxinus*.

Description. Conidiomata pycnidial, immersed in bark, scattered, slightly erumpent through the bark surface, nearly flat, discoid, with a single locule. Ectostromatic disc grey to dark brown, circular to ovoid, one ostiole per disc, 150–325

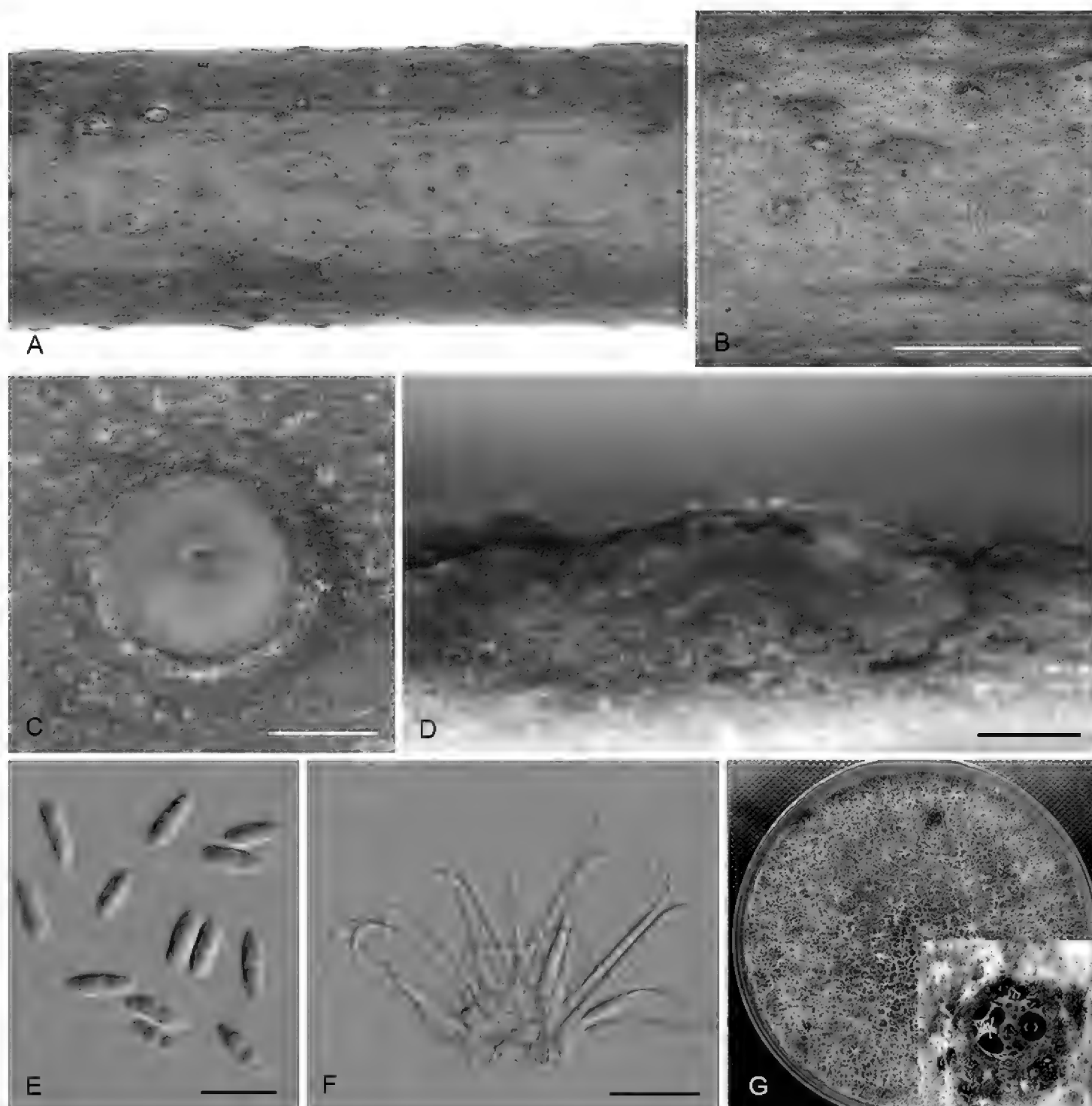


Figure 13. *Diaporthe fraxinicola* (CFCC 52582) **A–B** Habit of conidiomata on branches **C** Transverse section of conidioma **D** Longitudinal section of conidioma **E** Alpha conidia **F** Beta conidia **G** Culture on PDA and conidiomata. Scale bars: 500 μm (**B**), 200 μm (**C**), 100 μm (**D**), 10 μm (**E–F**).

μm diam. Locule circular, undivided, 275–480 μm diam. Conidiophores 10.5–17.5 \times 2.1–3.2 μm , hyaline, branched, cylindrical to clavate, straight, tapering towards the apex. Alpha conidia hyaline, aseptate, ellipsoidal to oval, 2–3-guttulate, rounded at both ends, 7–10 \times 2.9–3.2 μm (av. = 8.5 \times 3 μm , n = 30). Beta conidia hyaline, filiform, straight or hamate, eguttulate, aseptate, base subtruncate, tapering towards one apex, 19–29.5 \times 1.4 μm (av. = 24.5 \times 1.4 μm , n = 30).

Culture characters. Cultures incubated on PDA at 25 °C in darkness. Colony originally flat with white aerial mycelium, becoming yellowish, dense and felted aerial mycelium with age, with visible solitary or aggregated conidiomata at maturity.

Additional material examined. CHINA. Shaanxi Province: Zhashui city, Niubeiliang Reserve, on symptomatic twigs of *Fraxinus chinensis*, 7 July 2017, Q. Yang, living culture CFCC 52583 (BJFC-S1496).

Notes. This new species is introduced as molecular data, shows it to be a distinct clade with high support (ML/Bi=100/1) and it appears most closely related to *D. oraccinii* and *D. acerigena*. *Diaporthe fraxinicola* can be distinguished from *D. oraccinii* by 22 nucleotides in concatenated alignment, in which 6 were distinct in the ITS region, 8 in the *tef1* region and 8 in the *tub2* region; from *D. acerigena* by 27 nucleotides in concatenated alignment, in which 11 were distinct in the ITS region, 3 in the *tef1* region and 13 in the *tub2* region. Morphologically, *D. fraxinicola* differs from *D. oraccinii* in longer and larger alpha conidia (7–10 × 2.9–3.2 vs. 5.5–7.5 × 0.5–2 µm); differs from *D. acerigena* in larger alpha conidia (2.9–3.2 vs. 2.1–2.9 µm) (Gao et al. 2016).

***Diaporthe kadsurae* C.M. Tian & Q. Yang, sp. nov.**

MycoBank: MB824713

Figure 14

Diagnosis. *Diaporthe kadsurae* differs from its closest phylogenetic species *D. fusicola* and *D. ovoicicola* in ITS, *cal* and *tef1* loci based on the alignments deposited in TreeBASE.

Holotype. CHINA. Jiangxi Province: Shangrao city, Sanqing Mountain, on symptomatic branches of *Kadsura longipedunculata*, 1 Apr. 2017, B. Cao, Y.M. Liang & C.M. Tian (holotype: BJFC-S1497; ex-type culture: CFCC 52586).

Etymology. Named after the host genus on which it was collected, *Kadsura*.

Description. Conidiomata pycnidial, immersed in bark, scattered, slightly erumpent through the bark surface, nearly flat, discoid, with a single locule. Ectostromatic disc obviously, brown to black, one ostiole per disc. Locule undivided, 475–525 µm diam. Conidiophores 7–11 × 1.8–2.9 µm, cylindrical, hyaline, unbranched, straight or slightly curved, tapering towards the apex. Alpha conidia hyaline, aseptate, oval or fusoid, biguttulate, 5.5–7.5 × 2.1–2.9 µm (av. = 6.5 × 2.5 µm, n = 30). Beta conidia not observed.

Culture characters. Cultures incubated on PDA at 25 °C in darkness. Colony originally flat with white aerial mycelium, becoming dense and felted aerial mycelium in the centre and grey to black mycelium at the marginal area with solitary conidiomata at maturity.

Additional specimens examined. CHINA. Jiangxi Province: Shangrao city, Sanqing Mountain, on symptomatic branches of *Kadsura longipedunculata*, 1 Apr. 2017, B. Cao, Y.M. Liang & C.M. Tian, living culture CFCC 52587 (BJFC-S1498); Yunbifeng National Forest Park, on symptomatic twigs of *Acer* sp., 31 Mar. 2017, B. Cao, Y.M. Liang & C.M. Tian, living culture CFCC 52588 (BJFC-S1499); ibid. living culture CFCC 52589 (BJFC-S1500).

Notes. This new species is introduced as molecular data show it to be a distinct clade with high support (ML/Bi=100/1) and it appears most closely related to *D. fusi-*

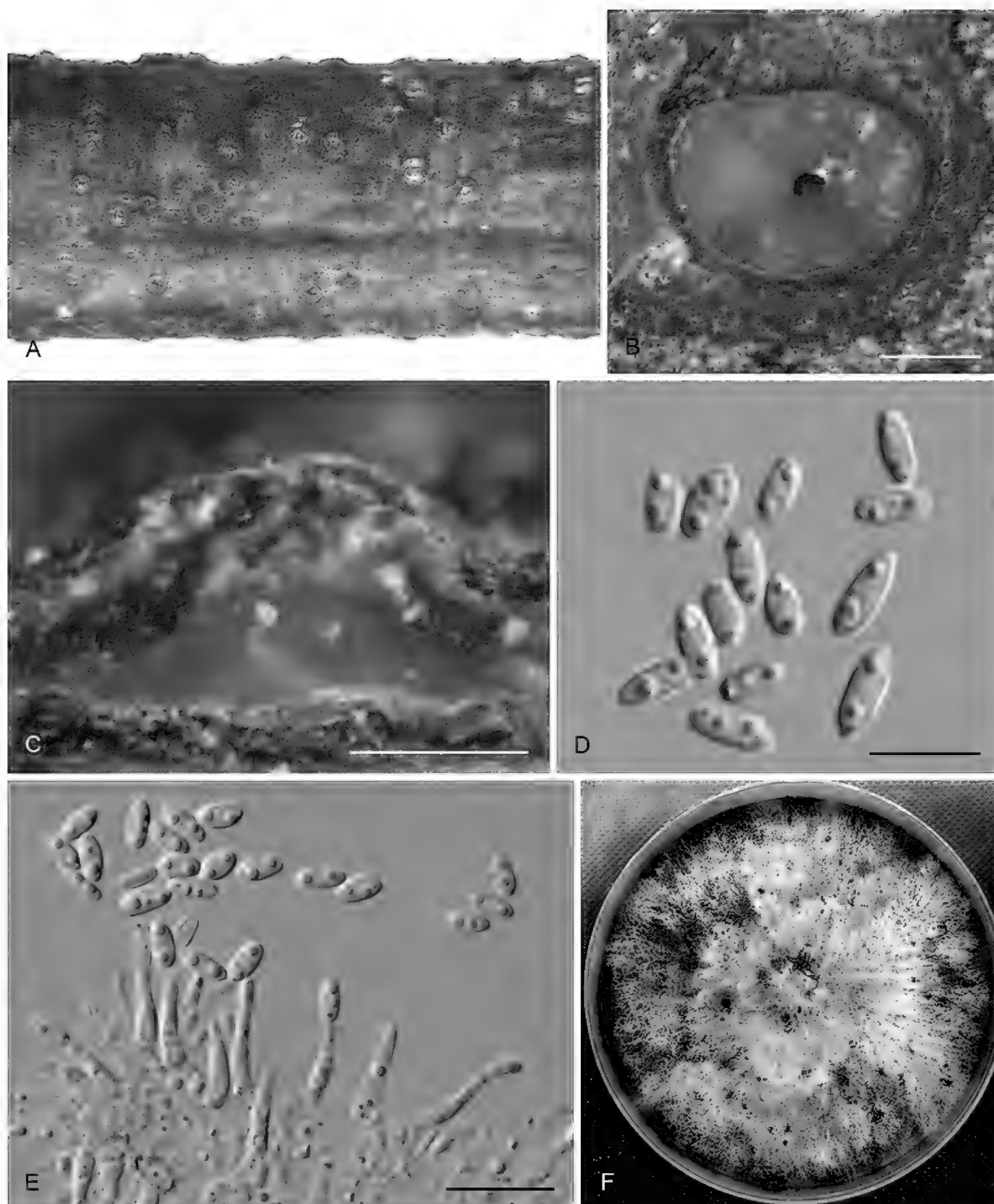


Figure 14. *Diaporthe kadsurae* (CFCC 52586) **A** Habit of conidiomata on branches **B** Transverse section of conidioma **C** Longitudinal section of conidioma **D** Alpha conidia **E** Conidiophores **F** Culture on PDA. Scale bars: 200 μ m (**B–C**), 10 μ m (**D–E**).

cola and *D. ovoicicola*. *Diaporthe kadsurae* can be distinguished from *D. fusicola* by 11 nucleotides in concatenated alignment, in which 4 were distinct in the ITS region and 7 in the *cal* region; from *D. ovoicicola* by 25 nucleotides in concatenated alignment, in which 12 were distinct in the ITS region, 6 in the *cal* region and 7 in the *tef1* region. Morphologically, *D. kadsurae* differs from *D. fusicola* and *D. ovoicicola* in shorter co-

nidiophores (7–11 μm in *D. kadsurae* vs. 11–24.1 μm in *D. fusicola*; 7–11 μm in *D. kadsurae* vs. 14.2–23.6 μm in *D. ovoicicola*) (Gao et al. 2014).

***Diaporthe padina* C.M. Tian & Q. Yang, sp. nov.**

MycoBank: MB824714

Figure 15

Diagnosis. *Diaporthe padina* can be distinguished from the phylogenetically closely related species *D. betulae* in smaller conidiomata and alpha conidia.

Holotype. CHINA. Heilongjiang Province: Liangshui Nature Reserve, on symptomatic twigs of *Padus racemosa*, 31 July 2016, Q. Yang (holotype: BJFC-S1501; ex-type culture: CFCC 52590).

Etymology. Named after the host genus on which it was collected, *Padus*.

Description. Conidiomata pycnidial, immersed in bark, scattered, slightly erumpent through the bark surface, discoid, with a single locule. Ectostromatic disc light brown, one ostiole per disc, 330–520 μm diam. Locule circular, undivided, 250–550 μm diam. Conidiophores 5.5–12.5 \times 1–1.5 μm , hyaline, unbranched, cylindrical, straight or slightly curved. Alpha conidia hyaline, aseptate, ellipsoidal to fusiform, eguttulate, 7–8 \times 1.5–2 μm (av. = 7.5 \times 1.8 μm , $n = 30$). Beta conidia hyaline, filiform, straight or hamate, eguttulate, aseptate, base truncate, 21–24 \times 1 μm (av. = 22 \times 1 μm , $n = 30$).

Culture characters. Cultures incubated on PDA at 25 °C in darkness. Colony originally flat with white aerial mycelium, becoming grey to brown in the centre, with pale grey, felted, valviform mycelium at the marginal area and aggregated conidiomata at maturity.

Additional material examined. CHINA. Heilongjiang Province: Liangshui Nature Reserve, on symptomatic twigs of *Padus racemosa*, 31 July 2016, Q. Yang, living culture CFCC 52591 (BJFC-S1502).

Notes. Four strains representing *D. padina* cluster in a well-supported clade and appear closely related to *D. betulae*. This species is phylogenetically closely related to, but clearly differentiated from, *D. betulae* by 40 different unique fixed alleles in ITS, *cal*, *his3*, *tef1* and *tub2* loci (4, 7, 10, 13 and 6 respectively) based on the alignments deposited in TreeBASE. Morphologically, *D. padina* differs from *D. betulae* in smaller conidiomata and alpha conidia (250–550 vs. 600–1250 μm in conidiomata; 7–8 \times 1.5–2 vs. 8.5–11 \times 3–4 μm in alpha conidia) (Du et al. 2016).

***Diaporthe ukurunduensis* C.M. Tian & Q. Yang, sp. nov.**

MycoBank: MB824715

Figure 16

Diagnosis. *Diaporthe ukurunduensis* can be distinguished from the phylogenetically closely related species *D. citrichinensis* in longer conidiophores and shorter alpha conidia.

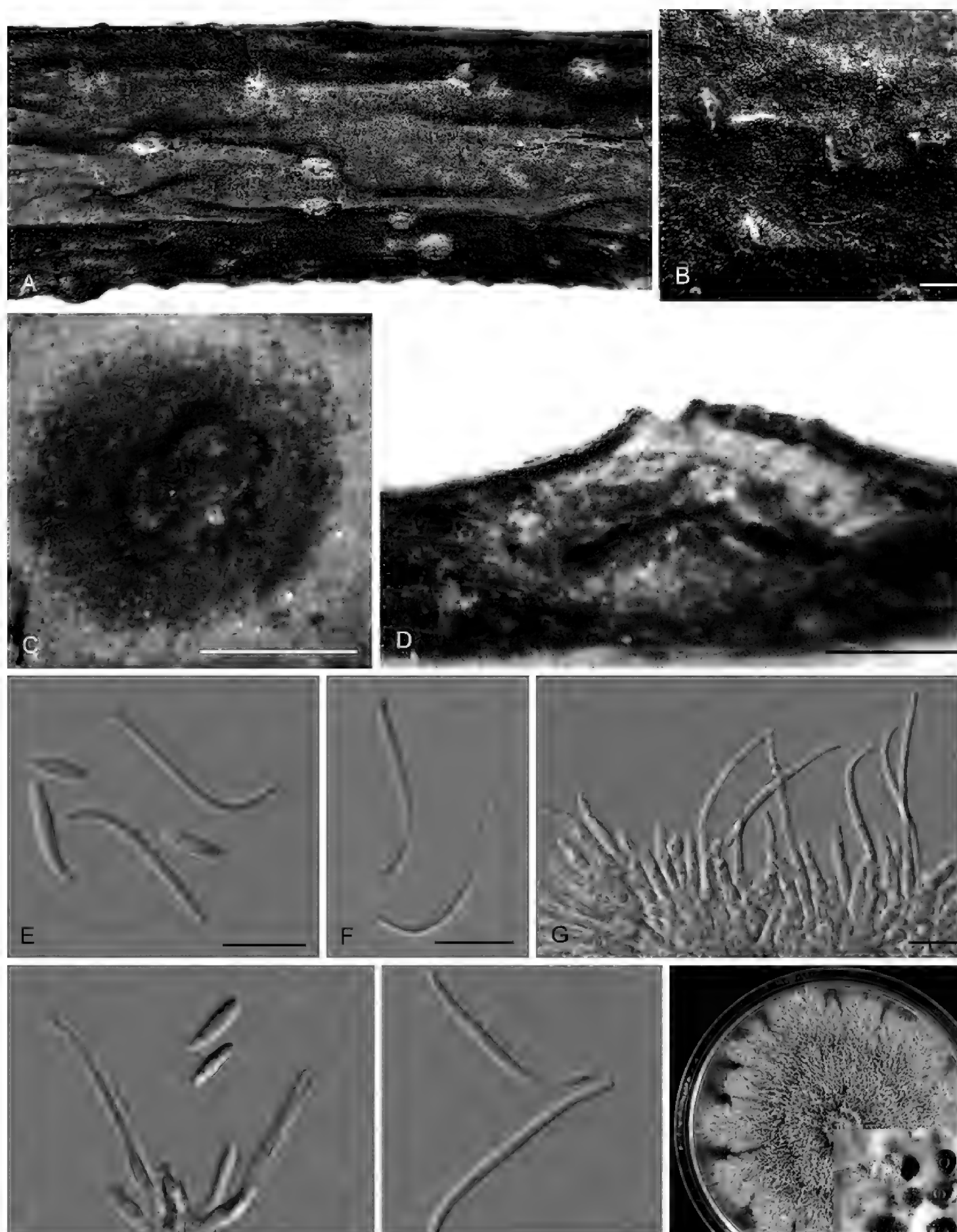


Figure 15. *Diaporthe padina* (CFCC 52590) **A–B** Habit of conidiomata on branches **C** Transverse section of conidioma **D** Longitudinal section of conidioma **E** Alpha and beta conidia **F, I** Beta conidia **G–H** Conidiophores **J** Culture on PDA and conidiomata. Scale bars: 500 μm (**B**), 200 μm (**C–D**), 10 μm (**E–I**).

Holotype. CHINA. Shaanxi Province: Qinling Mountain, on symptomatic twigs of *Acer ukurunduense*, 27 June 2017, Q. Yang (holotype: BJFC-S1503; ex-type culture: CFCC 52592).

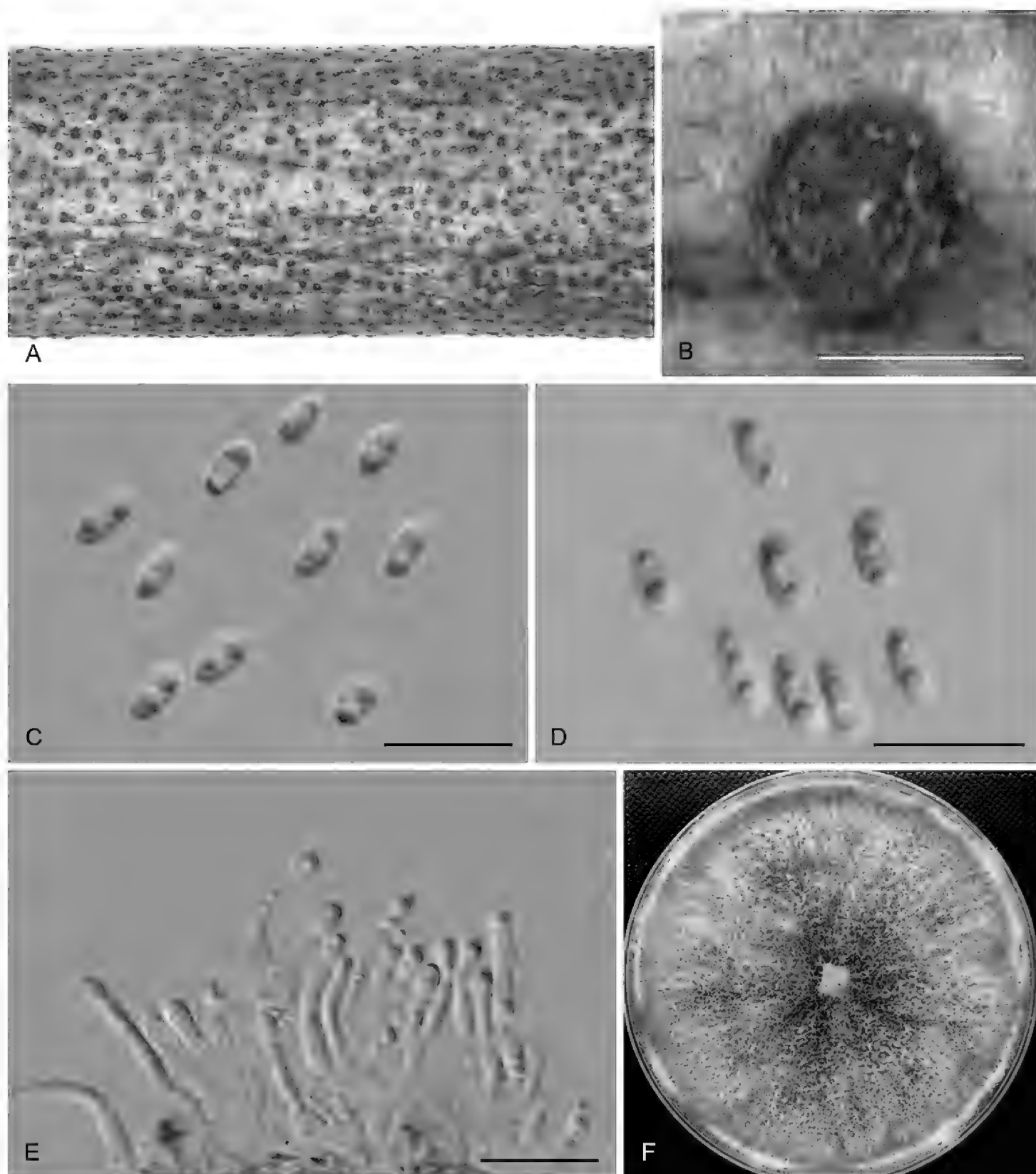


Figure 16. *Diaporthe ukurunduensis* (CFCC 52592) **A** Habit of conidiomata on branches **B** Transverse section of conidioma **C–D** Alpha conidia **E** Conidiophores **F** Culture on PDA. Scale bars: 200 μm (**B**), 10 μm (**C–E**).

Etymology. Named after the host species on which it was collected, *Acer ukurunduense*.

Description. Conidiomata pycnidial, immersed in bark, serried, slightly erumpent through the bark surface, nearly flat, discoid, with a single locule. Ectostromatic disc dark brown to black, one ostiole per disc. Locule circular, undivided, 165–215 μm diam. Conidiophores 11.5–18 \times 1.5 μm , hyaline, branched, cylindrical, straight or curved. Alpha conidia hyaline, aseptate, ellipsoidal to oval, biguttulate, 5–6 \times 2.1–2.9 μm (av. = 5.5 \times 2.5 μm , $n = 30$). Beta conidia not observed.

Culture characters. Cultures incubated on PDA at 25 °C in darkness. Colony originally flat with white aerial mycelium, becoming brown to pale black in the centre, dense, felted, conidiomata not observed.

Additional specimens examined. CHINA. Shaanxi Province: Qinling Mountain, on symptomatic twigs of *Acer ukurunduense*, 27 June 2017, Q. Yang, living culture CFCC 52593 (BJFC-S1503).

Notes. *Diaporthe ukurunduensis* comprises strains CFCC 52592 and CFCC 52593 closely related to *D. citrichinensis* in the combined phylogenetic tree (Fig. 1). *Diaporthe ukurunduensis* can be distinguished from *D. citrichinensis* based on ITS and *tef1* loci (10/470 in ITS and 4/336 in *tef1*).

Diaporthe unshiuensis F. Huang, K.D. Hyde & H.Y. Li, 2015

Figure 17

Description. On PNA: Conidiomata pycnidial, globose or rostrated, black, erumpent in tissue, erumpent at maturity, 260–500 µm diam, often with translucent conidial drops exuding from the ostioles. Conidiophores 18–28.5 × 1.4–2.1 µm, cylindrical, hyaline, branched, septate, straight or curved, tapering towards the apex. Alpha conidia abundant in culture, hyaline, aseptate, ellipsoidal to fusiform, biguttulate, sometimes with one end obtuse and the other acute, 6.5–8.5 × 2.1–2.5 µm (av. = 7.8 × 2.3 µm, n = 30). Beta conidia not observed.

Culture characters. Cultures incubated on PNA at 25 °C in darkness. Colony entirely white at surface, reverse with pale brown pigmentation, white, fluffy aerial mycelium.

Specimens examined. CHINA. Jiangsu Province: Nanjing city, on non-symptomatic twigs of *Carya illinoensis*, 10 Nov. 2015, Q. Yang, living culture CFCC 52594 and CFCC 52595 (BJFC-S1476).

Notes. *Diaporthe unshiuensis* was originally described from twigs of non-symptomatic *Fortunella margarita* in Zhejiang Province, China (Huang et al. 2015). In the present study, two isolates from twigs of asymptomatic *Carya illinoensis* were congruent with *D. unshiuensis* based on morphology and DNA sequences data (Fig. 1). We therefore describe *D. unshiuensis* as a known species for this clade.

Discussion

The current study described 15 *Diaporthe* species from 42 strains based on a large set of freshly collected specimens. It includes 12 new species and 3 known species, which were sampled from 16 host genera distributed over six Provinces of China (Table 1). In this study, 194 reference sequences (including outgroup) were selected based on BLAST searches of NCBI's GenBank nucleotide database and included in the phylogenetic analyses (Table 1). Phylogenetic analyses based on five combined loci (ITS, *cal*, *his3*, *tef1*

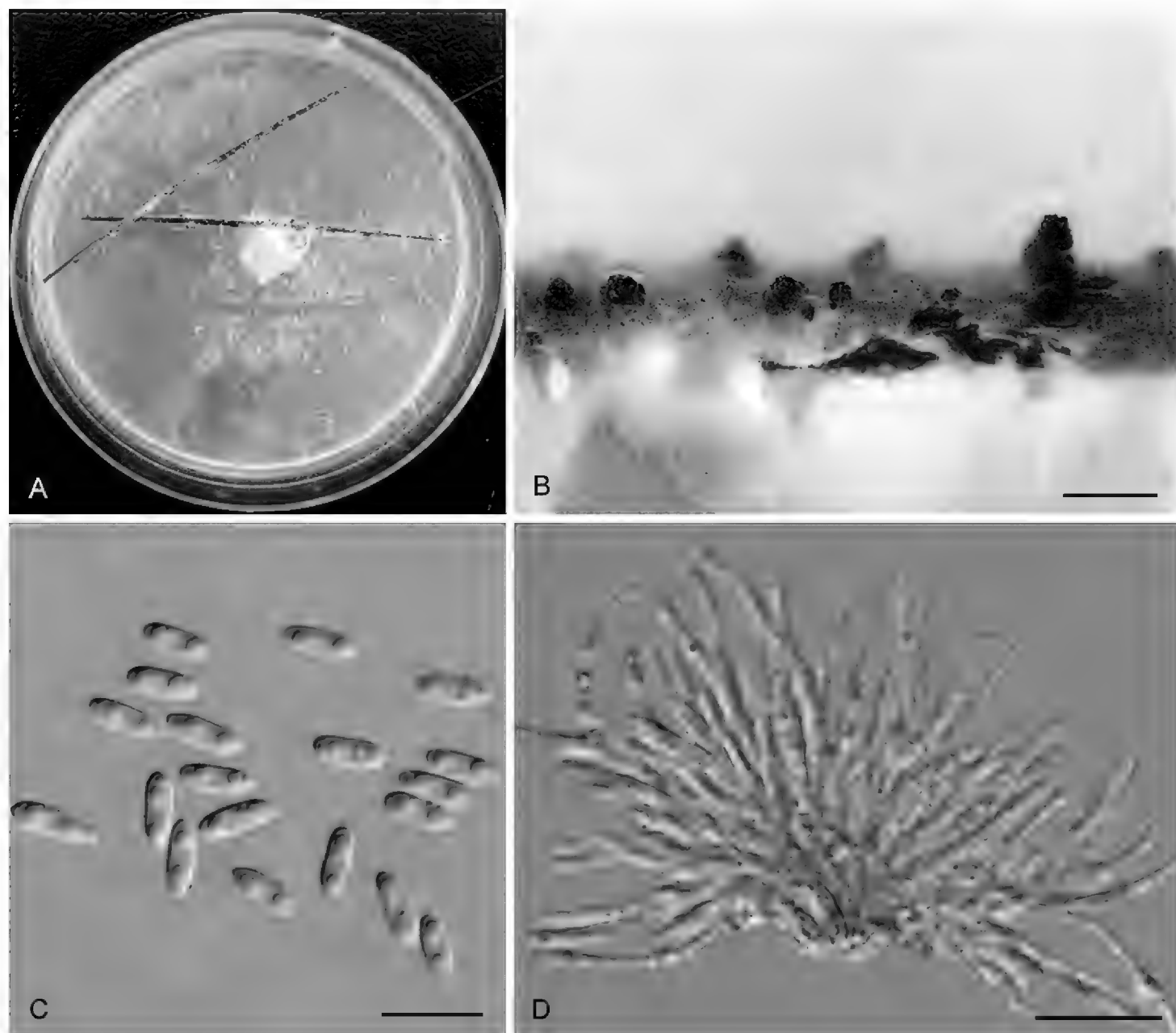


Figure 17. *Diaporthe unshiuensis* (CFCC 52594) **A** Culture on PNA **B** Conidiomata **C** Alpha conidia **D** Conidiophores. Scale bars: 500 μm (**B**), 10 μm (**C–D**).

and *tub2*), as well as morphological characters, revealed the diversity of *Diaporthe* species in China, mainly focusing on diebacks from major ecological or economic forest trees.

Several studies have been conducted associated with various hosts in China. For instance, the research conducted by Huang et al. (2015) revealed seven apparently undescribed endophytic *Diaporthe* species on *Citrus*. Gao et al. (2016) demonstrated that *Diaporthe* isolates, associated with *Camellia* spp., could be assigned to seven species and two species complexes. Recently, *Diaporthe* has been revealed as paraphyletic by Gao et al. (2017), showing that *Ophiodiaporthe*, *Pustulomyces*, *Phaeocystroma* and *Stenocarpella* embed in *Diaporthe* s. lat. and eight new species of *Diaporthe* were introduced from leaves of several hosts. However, the identification of *Diaporthe* species associated with dieback of forest trees has rarely been studied, thus a large-scale investigation of *Diaporthe* spp. was conducted from 2015 to 2017. This study provides the first molecular phylogenetic frame of *Diaporthe* diversity associated with dieback in China, combined with morphological descriptions.

Diaporthe eres, the type species of the genus, was initially described by Nitschke (1870), from *Ulmus* sp. collected in Germany. The major problem with this generic type was the lack of an ex-type culture or ex-epitype culture, although a broad species concept has historically been associated with *D. eres* (Udayanga et al. 2014b). Udayanga et al. (2014b) designed strain AR5193 as the epitype of *D. eres* and provided the phylogram of this complex using seven loci (ITS, *act*, *Apn2*, *cal*, *his3*, FG1093, *tef1* and *tub2*), amongst which the *tef1*, *Apn2* and *his3* genes were recognised as the best markers for defining species in the *D. eres* complex. Moreover, they showed that poorly supported non-monophyletic grouping was observed when ITS sequences were included in the combined analysis. In this study, although we conducted phylogenetic analysis as performed in previous studies on *Diaporthe* species (Santos et al. 2017), much confusion has, however, occurred in species separation of the *D. eres* complex (Fig. 1). Especially, the ITS region could lead to a confused taxonomic situation within this species complex. We found the three-gene analysis, excluding the ITS and *his3* regions, resulted in a more robust tree congruent with Udayanga et al. (2014b) and resolved the species boundaries within the *D. eres* species complex. The isolates, clustering with *D. eres* in this study, occur on multiple hosts from many different geographic locations. This study revealed three new species belonging to the *D. eres* complex, i.e. *D. betulina*, *D. chensiensis* and *D. padina*. It also shows *D. biguttusis*, *D. camptothecicola*, *D. ellipicola*, *D. longicicola*, *D. mahothocarpus* and *D. momicola* were clustered in *D. eres* and should be treated as synonyms of *D. eres*, which is in conformity with Fan et al. (2018).

The initial species concept of *Diaporthe*, based on the assumption of host-specificity, resulted in the introduction of more than 1000 taxa (<http://www.indexfungorum.org/>). Thus, during the past decade, a polyphasic approach, employing multi-locus DNA data together with morphology and ecology, has been employed for species boundaries in the genus (Crous et al. 2012, Udayanga et al. 2014a, b, Huang et al. 2015, Gao et al. 2016, 2017, Guarnaccia and Crous 2017, 2018, Hyde et al. 2017, 2018, Yang et al. 2017a, b, 2018, Guarnaccia et al. 2018, Jayawardena et al. 2018, Perera et al. 2018a, b, Tibpromma et al. 2018, Wanasinghe et al. 2018).

Further studies are required in order to conduct an extensive collection of *Diaporthe* isolates, to resolve taxonomic questions and to redefine species boundaries. Multiple strains from different locations should also be subjected to multi-gene phylogenetic analysis to determine intraspecific variation. The descriptions and molecular data of *Diaporthe* species provided in this study represent a resource for plant pathologists, plant quarantine officials and taxonomists for identification of *Diaporthe*.

Acknowledgements

This study is financed by National Natural Science Foundation of China (Project No.: 31670647). We are grateful to Chungen Piao, Minwei Guo (China Forestry Culture Collection Center (CFCC), Chinese Academy of Forestry, Beijing.

References

- Carbone I, Kohn LM (1999) A method for designing primer sets for speciation studies in filamentous ascomycetes. *Mycologia* 3: 553–556. <https://doi.org/10.2307/3761358>
- Chi PK, Jiang ZD, Xiang MM (2007) *Flora Fungorum Sinicorum*. Vol.34. *Phomopsis*. Science Press, Beijing.
- Crous PW, Gams W, Stalpers JA, Robert V, Stegehuis G (2004a) MycoBank: an online initiative to launch mycology into the 21st century. *Studies in Mycology* 50: 19–22.
- Crous PW, Groenewald JZ, Risède JM, Simoneau P, Hywel-Jones NL (2004b) *Calonectria* species and their *Cylindrocladium* anamorphs: species with sphaeropedunculate vesicles. *Studies in Mycology* 50: 415–430.
- Crous PW, Summerell BW, Shivas RG, Burgess TI, Decock CA, Dreyer LL, Granke LL, Guest DI, Hardy GEstJ, Hausbeck MK, Hüberli D, Jung T, Koukol O, Lennox CL, Liew ECY, Lombard L, McTaggart AR, Pryke JS, Roets F, Saude C, Shuttleworth LA, Stukely MJC, Vánky K, Webster BJ, Windstam ST, Groenewald JZ (2012) Fungal Planet description sheets: 107–127. *Persoonia: Molecular Phylogeny and Evolution of Fungi* 28: 138–182. <https://doi.org/10.3767/003158512X652633>
- Crous PW, Wingfield MJ, Le Roux JJ, Richardson DM, Strasberg D, Shivas RG, Alvarado P, Edwards J, Moreno G, Sharma R, Sonawane MS, Tan YP, Altés A, Barasubiye T, Barnes CM, Blanchette RA, Boertmann D, Bogo A, Carlavilla JR, Cheewangkoon R, Daniel R, de Beer ZW, de Jesús Yáñez-Morales M, Duong TA, Fernández-Vicente J, Geering ADW, Guest DI, Held BW, Heykoop M, Hubka V, Ismail AM, Kajale SC, Khemmuk W, Kolařík M, Kurli R, Lebeuf R, Lévesque CA, Lombard L, Magista D, Manjón JL, Marincowitz S, Mohedano JM, Nováková A, Oberlies NH, Otto EC, Paguigan ND, Pascoe IG, Pérez-Butrón JL, Perrone G, Rahi P, Raja HA, Rintoul T, Sanhueza RMV, Scarlett K, Shouche YS, Shuttleworth LA, Taylor PWJ, Thorn RG, Vawdrey LL, Solano-Vidal R, Voitek A, Wong PTW, Wood AR, Zamora JC, Groenewald JZ (2015) Fungal Planet description sheets: 371–399. *Persoonia: Molecular Phylogeny and Evolution of Fungi* 35: 264. <https://doi.org/10.3767/003158515X690269>
- Desjardins P, Hansen JB, Allen M (2009) Microvolume protein concentration determination using the NanoDrop 2000c spectrophotometer. *Journal of visualized experiments: JoVE* 33: 1–3. <https://doi.org/10.3791/1610>
- Diogo E, Santos JM, Phillips AJ (2010) Phylogeny, morphology and pathogenicity of *Diaporthe* and *Phomopsis* species on almond in Portugal. *Fungal Diversity* 44: 107–115. <https://doi.org/10.1007/s13225-010-0057-x>
- Dissanayake AJ, Phillips AJL, Hyde KD, Yan JY, Li XH (2017b) The current status of species in *Diaporthe*. *Mycosphere* 8(5): 1106–1156. <https://doi.org/10.5943/mycosphere/8/5/5>
- Dissanayake AJ, Zhang W, Liu M, Hyde KD, Zhao WS, Li XH, Yan JY (2017a) *Diaporthe* species associated with peach tree dieback in Hubei, China. *Mycosphere* 8: 533–549. <https://doi.org/10.5943/mycosphere/8/5/3>
- Doilom M, Dissanayake AJ, Wanasinghe DN, Wanasinghe DN, Boonmee S, Liu JK, Bhat DJ, Taylor JE, Bahkali AH, McKenzie EHC, Hyde KD (2017) Microfungi on *Tectona gran-*

- dis* (teak) in Northern Thailand. Fungal Diversity 82: 107–182. <https://doi.org/10.1007/s13225-016-0368-7>
- Doyle JJ, Doyle JL (1990) Isolation of plant DNA from fresh tissue. Focus 12: 13–15.
- Du Z, Fan XL, Hyde KD, Yang Q, Liang YM, Tian CM (2016) Phylogeny and morphology reveal two new species of *Diaporthe* from *Betula* spp. in China. Phytotaxa 269: 90–102. <https://doi.org/10.11646/phytotaxa.269.2.2>
- Erincik O, Madden L, Ferree D, Ellis M (2001) Effect of growth stage on susceptibility of grape berry and rachis tissues to infection by *Phomopsis viticola*. Plant Disease 85: 517–520. <https://doi.org/10.1094/PDIS.2001.85.5.517>
- Fan XL, Hyde KD, Udayanga D, Wu XY, Tian CM (2015) *Diaporthe rostrata*, a novel ascomycete from *Juglans mandshurica* associated with walnut dieback. Mycological Progress 14: 1–8. <https://doi.org/10.1007/s11557-015-1104-5>
- Fan XL, Yang Q, Bezerra JDP, Alvarez LV, Tian CM (2018) *Diaporthe* from walnut tree (*Juglans regia*) in China, with insight of *Diaporthe eres* complex. Mycological Progress: 1–13. <https://doi.org/10.1007/s11557-018-1395-4>
- Gao YH, Liu F, Cai L (2016) Unravelling *Diaporthe* species associated with *Camellia*. Systematics and Biodiversity 14: 102–117. <https://doi.org/10.1080/14772000.2015.1101027>
- Gao YH, Liu F, Duan W, Crous PW, Cai L (2017) *Diaporthe* is paraphyletic. IMA fungus 8: 153–187. <https://doi.org/10.5598/ima fungus.2017.08.01.11>
- Gao YH, Su YY, Sun W, Cai L (2015) *Diaporthe* species occurring on *Lithocarpus glabra* in China, with descriptions of five new species. Fungal Biology 115: 295–309. <https://doi.org/10.1016/j.funbio.2014.06.006>
- Gao YH, Sun W, Su YY, Cai L (2014) Three new species of *Phomopsis* in Gutianshan Nature Reserve in China. Mycological Progress 13: 111–121. <https://doi.org/10.1007/s11557-013-0898-2>
- Garcia-Reyne A, López-Medrano F, Morales JM, Esteban CG, Martín I, Eraña I, Meije Y, Lalueza A, Alastruey-Izquierdo A, Rodríguez-Tudela JL, Aguado JM (2011) Cutaneous infection by *Phomopsis longicolla* in a renal transplant recipient from Guinea: first report of human infection by this fungus. Transplant Infectious Disease 13: 204–207. <https://doi.org/10.1111/j.1399-3062.2010.00570.x>
- Glass NL, Donaldson GC (1995) Development of primer sets designed for use with the PCR to amplify conserved genes from filamentous ascomycetes. Applied and Environmental Microbiology 61: 1323–1330.
- Gomes RR, Glienke C, Videira SIR, Lombard L, Groenewald JZ, Crous PW (2013) *Diaporthe*: a genus of endophytic, saprobic and plant pathogenic fungi. Persoonia: Molecular Phylogeny and Evolution of Fungi 31: 1. <https://doi.org/10.3767/003158513X666844>
- Guarnaccia V, Crous PW (2017) Emerging citrus diseases in Europe caused by species of *Diaporthe*. IMA Fungus 8: 317–334. <https://doi.org/10.5598/ima fungus.2017.08.02.07>
- Guarnaccia V, Crous PW (2018) Species of *Diaporthe* on *Camellia* and *Citrus* in the Azores Islands. Phytopathologia Mediterranea. https://doi.org/10.14601/Phytopathol_Mediterr-23254
- Guarnaccia V, Groenewald JZ, Woodhall J, Armengol J, Cinelli T, Eichmeier A, Ezra D, Fontaine F, Gramaje D, Gutierrez-Aguirregabiria A, Kaliterna J, Kiss L, Larignon P, Luque J, Mugnai L, Naor V, Raposo R, Sándor E, Váczy KZ, Crous PW (2018) *Diaporthe* diversity

- and pathogenicity revealed from a broad survey of grapevine diseases in Europe. *Persoonia: Molecular Phylogeny and Evolution of Fungi* 40: 135–153. <https://doi.org/10.3767/persoonia.2018.40.06>
- Guarnaccia V, Vitale A, Cirvilleri G, Aiello D, Susca A, Epifani F, Perrone G, Polizzi G (2016) Characterisation and pathogenicity of fungal species associated with branch cankers and stem-end rot of avocado in Italy. *European Journal of Plant Pathology* 146: 963–976. <https://doi.org/10.1007/s10658-016-0973-z>
- Guindon S, Dufayard JF, Lefort V, Anisimova M, Hordijk W, Gascuel O (2010) New algorithms and methods to estimate maximum-likelihood phylogenies: assessing the performance of PhyML 3.0. *Systematic Biology* 59: 307–321. <https://doi.org/10.1093/sysbio/syq010>
- Hewitt W, Pearson R (1988) *Phomopsis* cane and leaf spot. *Compendium of grape diseases*: 17–18. APS Press, St Paul, Minnesota.
- Hillis DM, Bull JJ (1993) An empirical test of bootstrapping as a method for assessing confidence in phylogenetic analysis. *Systematic Biology* 42: 182–192. <https://doi.org/10.1093/sysbio/42.2.182>
- Huang F, Hou X, Dewdney MM, Fu Y, Chen G, Hyde KD, Li HY (2013) *Diaporthe* species occurring on *citrus* in China. *Fungal Diversity* 61: 237–250. <https://doi.org/10.1007/s13225-013-0245-6>
- Huang F, Udayanga D, Wang X, Hou X, Mei X, Fu Y, Hyde KD, Li HY (2015) Endophytic *Diaporthe* associated with *Citrus*: A phylogenetic reassessment with seven new species from China. *Fungal Biology* 119: 331–347. <https://doi.org/10.1016/j.funbio.2015.02.006>
- Hyde KD, Chaiwan N, Norphanphoun C, Boonmee S, Camporesi E, Chethana KWT, Dayarathne MC, de Silva NI, Dissanayake AJ, Ekanayaka AH, Hongsan S, Huang SK, Jayasiri SC, Jayawardena RS, Jiang HB, Karunarathna A, Lin CG, Liu JK, Liu NG, Lu YZ, Luo ZL, Maharachchimbura SSN, Manawasinghe IS, Pem D, Perera RH, Phukhamsakda C, Samarakoon MC, Senwanna C, Shang QJ, Tennakoon DS, Thambugala KM, Tibpromma S, Wanasinghe DN, Xiao YP, Yang J, Zeng XY, Zhang JF, Zhang SN, Bulgakov TS, Bhat DJ, Cheewangkoon R, Goh TK, Jones EBG, Kang JC, Jeewon R, Liu ZY, Lumyong S, Kuo CH, McKenzie EHC, Wen TC, Yan JY, Zhao Q (2018) *Mycosphere* notes 169–224. *Mycosphere* 9(2): 271–430. <https://doi.org/10.5943/mycosphere/9/2/8>
- Hyde KD, Norphanphoun C, Abreu VP, Bazzicalupo A, Chethana KT, Clericuzio M, Dayarathne MC, Dissanayake AJ, Ekanayaka AH, He MQ, Hongsan S, Huang SK, Jayasiri SC, Jayawardena RS, Karunarathna A, Konta S, Kušan I, Lee H, Li J, Lin CG, Liu NG, Lu YZ, Luo ZL, Manawasinghe IS, Mapook A, Perera RH, Phookamsak R, Phukhamsakda C, Siedlecki I, Soares AM, Tennakoon DS, Tian Q, Tibpromma S, Wanasinghe DN, Xiao YP, Yang J, Zeng XY, Abdel-Aziz FA, Li WJ, Senanayake IC, Shang QJ, Daranagama DA, de Silva NI, Thambugala KM, Abdel-Wahab MA, Bahkali AH, Berbee ML, Boonmee S, Bhat DJ, Bulgakov TS, Buyck B, Camporesi E, Castañeda-Ruiz RF, Chomnunti P, Doilom M, Dovana F, Gibertoni TB, Jadan M, Jeewon R, Jones EBG, Kang JC, Karunarathna SC, Lim YW, Liu JK, Liu ZY, Plautz Jr. HL, Lumyong S, Maharachchikumbura SSN, Matočec N, McKenzie EHC, Mešić A, Miller D, Pawłowska J, Pereira OL, Promptuttha I, Romero AI, Ryvarden L, Su HY, Suetrong S, Tkalc̆ec Z, Vizzini A, Wen TC, Wisitrassameewong K, Wrzosek M, Xu JC, Zhao Q, Zhao RL, Mortimer PE (2017) Fungal diversity

- notes 603–708: taxonomic and phylogenetic notes on genera and species. *Fungal Diversity* 87(1): 1–235. <https://doi.org/10.1007/s13225-017-0391-3>
- Jayawardena RS, Hyde KD, Chethana KWT, Daranagama DA, Dissanayake AJ, Goonasekara ID, Manawasinghe IS, Mapook A, Jayasiri SC, Karunarathna A, Li CG, Phukhamsakda C, Senanayake IC, Wanasinghe DN, Camporesi E, Bulgakov TS, Li XH, Liu M, Zhang W, Yan JY (2018) Mycosphere Notes 102–168: Saprotrophic fungi on *Vitis* in China, Italy, Russia and Thailand. *Mycosphere* 9(1): 1–114. <https://doi.org/10.5943/mycosphere/9/1/1>
- Katoh K, Toh H (2010) Parallelization of the MAFFT multiple sequence alignment program. *Bioinformatics* 26: 1899–1900. <https://doi.org/10.1093/bioinformatics/btq224>
- Lombard L, Van Leeuwen GCM, Guarnaccia V, Polizzi G, Van Rijswijk PC, Karin Rosendahl KC, Gabler J, Crous PW (2014) *Diaporthe* species associated with *Vaccinium*, with specific reference to Europe. *Phytopathologia Mediterranea* 53: 287–299. https://doi.org/10.14601/Phytopathol_Mediterr-14034
- Marin-Felix Y, Hernández-Restrepo M, Wingfield MJ, Akulov A, Carnegie AJ, Cheewangkoon R, Gramaje D, Groenewald JZ, Guarnaccia V, Halleen F, Lombard L, Luangsa-ard J, Marincowitz S, Moslemi A, Mostert L, Quaedvlieg W, Schumacher RK, Spies CFJ, Thangavel R, Taylor PWJ, Wilson AM, Wingfield BD, Wood AR, Crous PW (2018) Genera of phytopathogenic fungi: GOPHY2. *Studies in Mycology*. <https://doi.org/10.1016/j.simyco.2018.04.002>.
- Mostert L, Crous PW, Kang JC, Phillips AJ (2001) Species of *Phomopsis* and a *Libertella* sp. occurring on grapevines with specific reference to South Africa: morphological, cultural, molecular and pathological characterization. *Mycologia* 93: 146–167. <https://doi.org/10.2307/3761612>
- Mondal SN, Vincent A, Reis RF, Timmer LW (2007) Saprophytic colonization of citrus twigs by *Diaporthe citri* and factors affecting pycnidial production and conidial survival. *Plant Disease* 91: 387–392. <https://doi.org/10.1094/PDIS-91-4-0387>
- Muntañola-Cvetković M, Mihaljčević M, Petrov M (1981) On the identity of the causative agent of a serious *Phomopsis-Diaporthe* disease in sunflower plants. *Nova Hedwigia* 34: 417–435.
- Muralli TS, Suryanarayanan TS, Geeta R (2006) Endophytic *Phomopsis* species: host range and implications for diversity estimates. *Canadian Journal of Microbiology* 52: 673–680. <https://doi.org/10.1139/w06-020>
- Nitschke T (1870) *Pyrenomycetes Germanici* 2. Eduard Trewendt, Breslau, 245.
- O'Donnell K, Cigelnik E (1997) Two divergent intragenomic rDNA ITS2 types within a monophyletic lineage of the fungus *Fusarium* are nonorthologous. *Molecular Phylogenetics and Evolution* 7: 103–116. <https://doi.org/10.1006/mpev.1996.0376>
- Perera RH, Hyde KD, Dissanayake AJ, Jones EB, Liu JK, Wei D, Liu ZY (2018b) *Diaporthe collariana* sp. nov., with prominent collarettes associated with *Magnolia champaca* fruits in Thailand. *Studies in Fungi* 3(1): 141–151. <https://doi.org/10.5943/sif/3/1/16>
- Perera RH, Hyde KD, Peršoh D, Jones EBG, Liu JK, Liu ZY (2018a) Additions to wild seed and fruit fungi 1: The sexual morph of *Diaporthe rosae* on *Magnolia champaca* and *Senna siamea* fruits in Thailand. *Mycosphere* 9(2): 256–270. <https://doi.org/10.5943/mycosphere/9/2/7>

- Posada D, Crandall KA (1998) Modeltest: testing the model of DNA substitution. *Bioinformatics* 14: 817–818. <https://doi.org/10.1093/bioinformatics/14.9.817>
- Rannala B, Yang Z (1996) Probability distribution of molecular evolutionary trees: a new method of phylogenetic inference. *Journal of Molecular Evolution* 43: 304–311. <https://doi.org/10.1007/BF02338839>
- Rayner RW (1970) A mycological colour chart. Commonwealth Mycological Institute, Kew.
- Rehner SA, Uecker FA (1994) Nuclear ribosomal internal transcribed spacer phylogeny and host diversity in the coelomycete *Phomopsis*. *Canadian Journal of Botany* 72: 1666–1674. <https://doi.org/10.1139/b94-204>
- Rossmann AY, Adams GC, Cannon PF, Castlebury LA, Crous PW, Gryzenhout M, Jaklitsch WM, Mejia LC, Stoykov D, Udayanga D, Voglmayr H, Walker DM (2015) Recommendations of generic names in *Diaporthales* competing for protection or use. *IMA Fungus* 6: 145–154. <https://doi.org/10.5598/ima fungus.2015.06.01.09>
- Rossmann AY, Farr DE, Castlebury LA (2007) A review of the phylogeny and biology of the *Diaporthales*. *Mycoscience* 48: 135–144. <https://doi.org/10.1007/s10267-007-0347-7>
- Saccardo PA, Roumeguère C (1884) *Reliquiae mycologicae Libertianae*. IV. *Revue Mycologique*, Toulouse 6: 26–39.
- Santos JM, Correia VG, Phillips AJL (2010) Primers for mating-type diagnosis in *Diaporthe* and *Phomopsis*, their use in teleomorph induction in vitro and biological species definition. *Fungal Biology* 114: 255–270. <https://doi.org/10.1016/j.funbio.2010.01.007>
- Santos JM, Phillips AJL (2009) Resolving the complex of *Diaporthe* (*Phomopsis*) species occurring on *Foeniculum vulgare* in Portugal. *Fungal Diversity* 34: 111–125.
- Santos JM, Vrandečić K, Čosić J, Duvnjak T, Phillips AJL (2011) Resolving the *Diaporthe* species occurring on soybean in Croatia. *Persoonia: Molecular Phylogeny and Evolution of Fungi* 27: 9–19. <https://doi.org/10.3767/003158511X603719>
- Santos L, Alves A, Alves R (2017) Evaluating multi-locus phylogenies for species boundaries determination in the genus *Diaporthe*. *PeerJ* 5: e3120. <https://doi.org/10.7717/peerj.3120>
- Smith H, Wingfeld MJ, Coutinho TA, Crous PW (1996) *Sphaeropsis sapinea* and *Botryosphaeria dothidea* endophytic in *Pinus* spp. and *Eucalyptus* spp. in South Africa. *South African Journal of Botany* 62: 86–88. [https://doi.org/10.1016/S0254-6299\(15\)30596-2](https://doi.org/10.1016/S0254-6299(15)30596-2)
- Tai FL (1979) *Sylloge fungorum sinicorum*. Science press, 1527 pp.
- Tamura K, Stecher G, Peterson D, Filipowski A, Kumar S (2013) MEGA6: molecular evolutionary genetics analysis version 6.0. *Molecular Biology and Evolution* 30: 2725–2729. <https://doi.org/10.1093/molbev/mst197>
- Tanney JB, McMullin DR, Green BD, Miller JD, Seifert KA (2016) Production of antifungal and antiinsectan metabolites by the *Picea* endophyte *Diaporthe maritima* sp. nov. *Fungal Biology* 120: 1448–1457. <https://doi.org/10.1016/j.funbio.2016.05.007>
- Teng SC (1963) *Fungi of China*. Science press, 808 pp.
- Thompson SM, Tan YP, Shivas RG, Neate SM, Morin L, Bissett A, Aitken EAB (2015) Green and brown bridges between weeds and crops reveal novel *Diaporthe* species in Australia. *Persoonia: Molecular Phylogeny and Evolution of Fungi* 35: 39. <https://doi.org/10.3767/003158515X687506>

- Thompson SM, Tan YP, Young AJ, Neate SM, Aitken EAB, Shivas RG (2011) Stem cankers on sunflower (*Helianthus annuus*) in Australia reveal a complex of pathogenic *Diaporthe* (*Phomopsis*) species. *Persoonia: Molecular Phylogeny and Evolution of Fungi* 27: 80. <https://doi.org/10.3767/003158511X617110>
- Tibpromma S, Hyde KD, Bhat JD, Mortimer PE, Xu J, Prompttha I, Mingkwan D, Yang JB, Tang AMC, Karunarathna SC (2018) Identification of endophytic fungi from leaves of Pandanaceae based on their morphotypes and DNA sequence data from southern Thailand. *MycoKeys* 33: 25. <https://doi.org/10.3897/mycokeys.33.23670>
- Udayanga D, Castlebury LA, Rossman AY, Chukeatirote E, Hyde KD (2014b) Insights into the genus *Diaporthe*: phylogenetic species delimitation in the *D. eres* species complex. *Fungal Diversity* 67: 203–229. <https://doi.org/10.1007/s13225-014-0297-2>
- Udayanga D, Castlebury LA, Rossman AY, Chukeatirote E, Hyde KD (2015) The *Diaporthe sojae* species complex: Phylogenetic re-assessment of pathogens associated with soybean, cucurbits and other field crops. *Fungal Biology* 119: 383–407. <https://doi.org/10.1016/j.funbio.2014.10.009>
- Udayanga D, Castlebury LA, Rossman AY, Hyde KD (2014a) Species limits in *Diaporthe*: molecular re-assessment of *D. citri*, *D. cytosporella*, *D. foeniculina* and *D. rudis*. *Persoonia: Molecular Phylogeny and Evolution of Fungi* 32: 83–101. <https://doi.org/10.3767/003158514X679984>
- Udayanga D, Liu X, Crous PW, McKenzie EH, Chukeatirote E, Chukeatirote E, Hyde KD (2012a) A multi-locus phylogenetic evaluation of *Diaporthe* (*Phomopsis*). *Fungal Diversity* 56: 157–171. <https://doi.org/10.1007/s13225-012-0190-9>
- Udayanga D, Liu X, McKenzie EH, Chukeatirote E, Bahkali AH, Hyde KD (2011) The genus *Phomopsis*: biology, applications, species concepts and names of common phytopathogens. *Fungal Diversity* 50: 189–225. <https://doi.org/10.1007/s13225-011-0126-9>
- Udayanga D, Liu X, McKenzie EH, Chukeatirote E, Hyde KD (2012b) Multi-locus phylogeny reveals three new species of *Diaporthe* from Thailand. *Cryptogamie, Mycologie* 33: 295–309. <https://doi.org/10.7872/crym.v33.iss3.2012.295>
- Uecker FA (1988) A world list of *Phomopsis* names with notes on nomenclature, morphology and biology. *Mycological Memoirs* 13: 1–231.
- Wanasinghe DN, Phukhamsakda C, Hyde KD, Jeewon R, Lee HB, Jones EG, Tibpromma S, Tennakoon DS, Dissanayake AJ, Jayasiri SC, Gafforov Y, Camporesi E, Bulgakov TS, Ekanayake AH, Perera RH, Samarakoon MC, Goonasekara ID, Mapook A, Li WJ, Senanayake IC, Li J, Norphanphoun C, Doilom M, Bahkali AH, Xu JC, Mortimer PE, Tibbell L, Tibell S, Karunarathna SC (2018) Fungal diversity notes 709–839: taxonomic and phylogenetic contributions to fungal taxa with an emphasis on fungi on Rosaceae. *Fungal Diversity* 89(1): 1–236. <https://doi.org/10.1007/s13225-018-0395-7>
- Wei JC (1979) Identification of Fungus Handbook. Science press, 780 pp.
- White TJ, Bruns T, Lee S, Taylor J (1990) Amplification and direct sequencing of fungal ribosomal RNA genes for phylogenetics. *PCR Protocols: A Guide to Methods and Applications* 18: 315–322.

- Yang Q, Du Z, Tian CM (2018) Phylogeny and morphology reveal two new species of *Diaporthe* from Traditional Chinese Medicine in Northeast China. *Phytotaxa* 336: 159–170. <https://doi.org/10.11646/phytotaxa.336.2.3>
- Yang Q, Fan XL, Du Z, Liang YM, Tian CM (2017c). *Diaporthe camptothecicola* sp. nov. on *Camptotheca acuminata* in China. *Mycotaxon* 132: 591–601. <https://doi.org/10.5248/132.591>
- Yang Q, Fan XL, Du Z, Tian CM (2017a) *Diaporthe* species occurring on *Senna bicapsularis* in southern China, with descriptions of two new species. *Phytotaxa* 302: 145–155. <https://doi.org/10.11646/phytotaxa.302.2.4>
- Yang Q, Fan XL, Du Z, Tian CM (2017b) *Diaporthe juglandicola* sp. nov. (Diaporthales, Ascomycetes), evidenced by morphological characters and phylogenetic analysis. *Mycosphere* 8: 817–826. <https://doi.org/10.5943/mycosphere/8/5/3>